Q2

PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

To:

Assistant Commissioner for Patents
United States Patent and Trademark Office
Box PCT
Washington, D.C.20231
ÉTATS-UNIS D'AMÉRIQUE

in its capacity as elected Office

Date of mailing (day/month/year) 28 February 2000 (28.02.00)	
International application No. PCT/GB99/02123	**Applicant's or agent's file reference** AHB/CP5775333
International filing date (day/month/year) 02 July 1999 (02.07.99)	**Priority date** (day/month/year) 02 July 1998 (02.07.98)
Applicant WILLIAMS, Andrew, James et al	

1. The designated Office is hereby notified of its election made:

[X] in the demand filed with the International Preliminary Examining Authority on:

01 February 2000 (01.02.00)

[ ] in a notice effecting later election filed with the International Bureau on:

2. The election [X] was

[ ] was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO **34, chemin des Colombettes** **1211 Geneva 20, Switzerland**	Authorized officer Juan Cruz
Facsimile No.: (41-22) 740.14.35	Telephone No.: (41-22) 338.83.38

Form PCT/IB/331 (July 1992)

3132768

PCT

REQUEST

The undersigned requests that the present international application be processed according to the Patent Cooperation Treaty

For receiving Office use only

International Application No.

International Filing Date

Name of receiving Office and "PCT International Application"

Applicant's or agent's file reference *(if desired) (12 characters maximum)* AHB/CP5775333

Box No. I TITLE OF INVENTION SPECIFIC BINDING PROTEINS INCLUDING ANTIBODIES WHICH BIND TO THE NECROTIC CENTRE OF TUMOURS, AND USES THEREOF

Box No. II APPLICANT

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.)*

CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED
THE SCIENCE PARK
MELBOURN
ROYSTON
CAMBRIDGESHIRE SG8 6JJ
UNITED KINGDOM

☐ This person is also inventor.

Telephone No.

Facsimile No.

Teleprinter No.

State *(that is, country) of nationality:* GB

State *(that is, country) of residence:* GB

This person is applicant for the purposes of: ☐ all designated States ☒ all designated States except the United States of America ☐ the United States of America only ☐ the States indicated in the Supplemental Box

Box No. III FURTHER APPLICANT(S) AND/OR (FURTHER) INVENTOR(S)

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.)*

WILLIAMS Andrew James
LARKSMEAD
LONG LANE
FOWLMERE
ROYSTON SG8 7TG
UNITED KINGDOM

This person is:

☐ applicant only

☒ applicant and inventor

☐ inventor only *(if this check-box is marked, do not fill in below.)*

State *(that is, country) of nationality:* GB

State *(that is, country) of residence:* GB

This person is applicant for the purposes of: ☐ all designated ☐ all designated States except the United States of America ☒ the United States of America only ☐ the States indicated in the Supplemental Box

☒ Further applicants and/or (further) inventors are indicated on a continuation sheet.

Box No. IV AGENT OR COMMON REPRESENTATIVE; OR ADDRESS FOR CORRESPONDENCE

The person identified below is hereby/has been appointed to act on behalf of the applicant(s) before the competent International Authorities as: ☒ agent ☐ common representative

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country.)*

BRASNETT, ADRIAN H. and others
MEWBURN ELLIS
YORK HOUSE
23 KINGSWAY
LONDON WC2B 6HP
GB

Telephone No. 0171 240 4405

Facsimile No. +44 171 240 9339

Teleprinter No.

☐ Mark this check-box where no agent or common representative is/has been appointed and the space above is used instead to indicate a special address to which correspondence should be sent.

Continuation of Box No. III FURTHER APPLICANTS AND/OR (FURTHER) INVENTORS

If none of the following sub-boxes is used, this sheet is not to be included in the request.

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.)*

TEMPEST Philip Ronald
43 HIGH STREET
WEST WRATTING
CAMBRIDGESHIRE CB1 5LU
UNITED KINGDOM

This person is:
- ☐ applicant only
- ☒ applicant and inventor
- ☐ inventor only *(if this check-box is marked, do not fill in below.)*

State *(that is, country) of nationality:* GB

State *(that is, country) of residence:* GB

This person is applicant for the purposes of:
- ☐ all designated states
- ☐ all designated States except the United States of America
- ☒ the United States of America only
- ☐ the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.))*

HOLTET Thor Las
MORKHOJ BYGADE 16
2860 SOBORG
DENMARK

This person is:
- ☐ applicant only
- ☒ applicant and inventor
- ☐ inventor only *(if this check-box is marked, do not fill in below.)*

State *(that is, country) of nationality:* DK

State *(that is, country) of residence:* DK

This person is applicant for the purposes of:
- ☐ all designated states
- ☐ all designated States except the United States of America
- ☒ the United States of America only
- ☐ the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.))*

JACKSON Helen
PEACE COTTAGE
15 BEDFORD ROAD
NORTH HILL
BEDFORDSHIRE SG18 9AH
UNITED KINGDOM

This person is:
- ☐ applicant only
- ☒ applicant and inventor
- ☐ inventor only *(if this check-box is marked, do not fill in below.)*

State *(that is, country) of nationality:* GB

State *(that is, country) of residence:* GB

This person is applicant for the purposes of:
- ☐ all designated states
- ☐ all designated States except the United States of America
- ☒ the United States of America only
- ☐ the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (that is, country) of residence if no State of residence is indicated below.))*

This person is:
- ☐ applicant only
- ☐ applicant and inventor
- ☐ inventor only *(if this check-box is marked, do not fill in below.)*

State *(that is, country) of nationality:*

State *(that is, country) of residence:*

This person is applicant for the purposes of:
- ☐ all designated states
- ☐ all designated States except the United States of America
- ☐ the United States of America only
- ☐ the States indicated in the Supplemental Box

☐ Further applicants and/or (further) inventors are indicated on another continuation sheet.

Box No. V DESIGNATION OF STATES

The following designations are hereby made under Rule 4.9(a) *(mark the applicable check-boxes; at least one must be marked)*:

Regional Patent

- [X] **AP** — **ARIPO Patent:** GH Ghana, GM Gambia, KE Kenya, LS Lesotho, MW Malawi, SD Sudan, SL Sierra Leone, SZ Swaziland, UG Uganda, ZW Zimbabwe, and any other State which is a Contracting State of the Harare Protocol and of the PCT
- [X] **EA** — **Eurasian Patent:** AM Armenia, AZ Azerbaijan, BY Belarus, KG Kyrgyzstan, KZ Kazakstan, MD Republic of Moldova, RU Russian Federation, TJ Tajikistan, TM Turkmenistan, and any other State which is a Contracting State of the Eurasian Patent Convention and of the PCT
- [X] **EP** — **European Patent:** AT Austria, BE Belgium, CH and LI Switzerland and Liechtenstein, CY Cyprus, DE Germany, DK Denmark, ES Spain, FI Finland, FR France, GB United Kingdom, GR Greece, IE Ireland, IT Italy, LU Luxembourg, MC Monaco, NL Netherlands, PT Portugal, SE Sweden, and any other State which is a Contracting State of the European Patent Convention and of the PCT
- [X] **OA** — **OAPI Patent:** BF Burkina Faso, BJ Benin, CF Central African Republic, CG Congo, CI Côte d'Ivoire, CM Cameroon, GA Gabon, GN Guinea, GW Guinea-Bissau, ML Mali, MR Mauritania, NE Niger, SN Senegal, TD Chad, TG Togo, and any other State which is a member State of OAPI and a Contracting State of the PCT *(if other kind of protection or treatment desired, specify on dotted line)*

.....

National Patent *(if other kind of protection desired, specify on dotted line)*:

- [X] AE United Arab Emirates
- [X] AL Albania.....
- [X] AM Armenia.....
- [X] AT Austria.....
- [X] AU Australia.....
- [X] AZ Azerbaijan
- [X] BA Bosnia & Herzegovina.....
- [X] BB Barbados
- [X] BG Bulgaria.....
- [X] BR Brazil.....
- [X] BY Belarus.....
- [X] CA Canada
- [X] CH and LI Switzerland and Liechtenstein.....
- [X] CN China.....
- [X] CU Cuba.....
- [X] CZ Czech Republic.....
- [X] DE Germany.....
- [X] DK Denmark.....
- [X] EE Estonia.....
- [X] ES Spain.....
- [X] FI Finland.....
- [X] GB United Kingdom.
- [X] GD Grenada
- [X] GE Georgia.....
- [X] GH Ghana.....
- [X] GM Gambia
- [X] HR Croatia.....
- [X] HU Hungary .....
- [X] ID Indonesia
- [X] IL Israel.....
- [X] IN India
- [X] IS Iceland
- [X] JP Japan.....
- [X] KE Kenya.....
- [X] KG Kyrgyzstan.....
- [X] KP Democratic People's Republic of Korea.....
- [X] KR Republic of Korea.....
- [X] KZ Kazakstan .....
- [X] LC St Lucia
- [X] LK Sri Lanka
- [X] LR Liberia.
- [X] LS Lesotho.....
- [X] LT Lithuania
- [X] LU Luxembourg
- [X] LV Latvia
- [X] MD Republic of Moldova .....
- [X] MG Madagascar .....
- [X] MK The former Yugoslav Republic of Macedonia...
- [X] MN Mongolia .....
- [X] MW Malawi.....
- [X] MX Mexico.....
- [X] NO Norway
- [X] NZ New Zealand .....
- [X] PL Poland .....
- [X] PT Portugal .....
- [X] RO Romania
- [X] RU Russian Federation .....
- [X] SD Sudan
- [X] SE Sweden
- [X] SG Singapore
- [X] SI Slovenia .....
- [X] SK Slovakia .....
- [X] SL Sierra Leone
- [X] TJ Tajikistan .....
- [X] TM Turkmenistan .....
- [X] TR Turkey .....
- [X] TT Trinidad and Tobago .....
- [X] UA Ukraine .....
- [X] UG Uganda .....
- [X] US United States of America.....
- [X] UZ Uzbekistan .....
- [X] VN Viet Nam .....
- [X] YU Yugoslavia .....
- [X] ZA South Africa
- [X] ZW Zimbabwe .....

Check-boxes reserved for designating States (for the purposes of a national patent) which have become party to the PCT after issuance of this sheet:

- [X]
- [X]
- [X] Any other state which is party to the PCT .....

Precautionary Designation Statement: In addition to the designations made above, the applicant also makes under Rule 4.9(b) all designations which would be permitted under the PCT except any designation(s) indicated in the Supplemental Box as being excluded from the scope of this statement.
The applicant declares that those additional designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be regarded as withdrawn by the applicant at the expiration of that time limit. *(Confirmation of a designation consists of the filing of a notice specifying that designation and the payment of the designation and confirmation fees. Confirmation must reach the receiving Office within the 15-month time limit.)*

Supplemental B x ***If the Supplemental Box is not used, this sheet need not be included in the request.***

Use this box in the following cases:

1. If, in any of the Boxes, the space is insufficient to furnish all the information:	***In such case, write "Continuation of Box No. ..." (indicate the number of the Box) and furnish the information in the same manner as required according to the captions of the Box in which the space was insufficient;***
in particular:	
(i) if more than two persons are involved as applicants and/or inventors and no "continuation sheet" is available:	*in such case, write "Continuation of Box III" and indicate for each additional person the same type of information as required in Box No. III. The country of the address indicated in this box is the applicant's state (that is, country) of residence if no state of residence is indicated below;*
(ii) if, in Box No. II or in any of the sub-boxes of Box No. III, the indication "the States indicated in the Supplemental Box" is checked:	*in such case, write "Continuation of Box No. II" or "Continuation of Box No. III" or "Continuation of Boxes No. II and No. III" (as the case may be), indicate the name of the applicant(s) involved and next to (each) such name, the State(s) (and/or, where applicable, ARIPO, Eurasian, European or OAPI patent) for the purposes of which the named person is applicant;*
(iii) if, in Box No. II or in any of the sub-boxes of Box No. III, the inventor or the inventor/applicant is not inventor for the purposes of all designated States or for the purposes of the United States of America:	*in such case, write "Continuation of Box No. II" or "Continuation of Box No. III" or "Continuation of Boxes No. II and No. III" (as the case may be), indicate the name of the inventor(s) and, next to (each) such name, the State(s) (and/or, where applicable, ARIPO, Eurasian, European or OAPI patent) for the purposes of which the named person is inventor;*
(iv) if, in addition to the agent(s) indicated in Box No. IV, there are further agents:	*in such case, write "Continuation of Box No. IV" and indicate for each further agent the same type of information as required in Box No. IV;*
(v) if, in Box No. V, the name of any State (or OAPI) is accompanied by the indication "patent of addition," or "certificate of addition," or if, in Box No. V, the name of the United States of America is accompanied by an indication "Continuation" or "Continuation-in-part":	*in such case, write "Continuation of Box No. V" and the name of each State involved (or OAPI), and after the name of each such State (or OAPI), the number of the parent title or parent application and the date of grant of the parent title or filing of the parent application;*
(vi) if, in Box No. VI, there are more than three earlier applications whose priority is claimed:	*in such case, write "Continuation of Box No. VI" and indicate for each additional earlier application the same type of information as required in Box No. VI.*
(vii) if, in Box No. VI, the earlier application is an ARIPO application:	*in such case, write "Continuation of Box No. VI", specify the number of the item corresponding to that earlier application and indicate at least one country party to the Paris Convention for the Protection of Industrial Property for which that earlier application was filed.*
2. If, with regard to the precautionary designation statement contained in Box No. V, the applicant wishes to exclude any State(s) from the scope of that statement:	***in such case, write "Designation(s) excluded from precautionary designation statement" and indicate the name or two-letter code of each state so excluded.***
3. If the applicant claims, in respect of any designated Office, the benefits of provisions of the national law concerning non-prejudicial disclosures or exceptions to lack of novelty:	***in such case, write "Statement Concerning Non-Prejudicial Disclosures or Exceptions to Lack of Novelty" and furnish that statement below.***

Continuation of Box IV

ARMITAGE, IAN M.
BRASNETT, ADRIAN H.
BREWSTER, ANDREA R.
CALDERBANK, T. ROGER
CARTER, STEPHEN
COLEIRO, RAYMOND
CRIPPS, JOANNA E
FORD, MICHAEL F.
GURA, H. ALAN
HACKNEY, NIGEL J.
HARRISON, DAVID C.
KIDDLE, SIMON J.
KREMER, SIMON M.
LINN, S. JONATHAN
LYONS, JUNE, M.
NICHOLLS, KATHRYN M.
O'BRIEN, CAROLINE J.
PAGET, HUGH C.E.
PAGET, HUGH C.E.
SANDERSON, MICHAEL J.
STONER, G. PATRICK
STUART, IAN
WALTON, SEÁN M.
WATKIN, TIMOTHY L.

Continuation of Box No. VIII

With regard to the formal sequence listing filed herewith, I confirm that the information recorded on the diskette is identical to the paper sequence listing.

B x No. VI **PRIORITY CLAIM** ☐ Further priority claims are indicated in the Supplemental Box

Filing date of earlier application (day/month/year)	Number of earlier application	Where earlier application is: national application: country	regional application:* regional Office	international application: receiving Office
item (1) 2 July 1998	9814383.7	GB		
item (2)				
item (3)				

☒ The receiving Office is requested to prepare and transmit to the International Bureau a certified copy of the earlier application(s) *(only if the earlier application was filed with the Office which for the purposes of the present international application is the receiving Office)* identified above as item(s): (1)

* *Where the earlier application is an ARIPO application, it is mandatory to indicate in the supplemental box at least one country party to the Paris Convention for the Protection of Industrial Property for which that earlier application was filed (Rule 4.10(b)(ii)). See Supplemental Box.*

Box No. VII **INTERNATIONAL SEARCHING AUTHORITY**

Choice of International Searching Authority (ISA) *(If two or more International Searching Authorities are competent to carry out the international search, indicate the Authority chosen; the two-letter code may be used):*

ISA /

Request to use results of earlier search; reference to that search (if an earlier search has been carried out by or requested from the International Searching Authority):

Date *(day/month/year)* Number Country *(or regional Office)*

Box No. VIII **CHECK LIST; LANGUAGE OF FILING**

This international application contains the following number of sheets	
request	:5
description (excluding sequence listing part)	:41
claims	:3
abstract	:1
drawings	:8
sequence listing part of description	:10
Total number of sheets	:68

This international application is **accompanied by** the item(s) marked below:

1. ☒ fee calculation sheet
2. ☐ separate signed power of attorney
3. ☒ copy of general power of attorney; reference number, if any:
4. ☐ statement explaining lack of signature
5. ☒ priority document(s) *identified in Box No. VI as item*(s):
6. ☐ translation of international application into (language):
7. ☐ separate indications concerning deposited microorganisms or other biological matter
8. ☒ nucleotide and/or amino acid sequence listing in computer readable form
9. ☒ other *(specify)*:Form 23/77 (x1)

Figure of the drawings which should accompany the abstract: 0

Language of filing of the international application: ENGLISH

Box No. IX **SIGNATURE OF APPLICANT OR AGENT**

Next to each signature indicate the name of the person signing and the capacity in which the person signs (if such capacity is not obvious from reading the request).

.....

HARRISON, DAVID C.
APPOINTED AGENT

For receiving Office use only

1. Date of actual receipt of the purported international application:	2. Drawings:
3. Corrected date of actual receipt due to later but timely received papers or drawings completing the purported international application:	☐ received:
4. Date of timely receipt of the required corrections under PCT Article 11(2):	☐ not received:
5. International Searching Authority (if two or more are competent): ISA/	6. ☐ Transmittal of search copy delayed until search fee is paid

For International Bureau use only

Date of receipt of the record copy by the International Bureau:

From the INTERNATIONAL BUREAU

PCT

NOTICE INFORMING THE APPLICANT OF THE COMMUNICATION OF THE INTERNATIONAL APPLICATION TO THE DESIGNATED OFFICES

(PCT Rule 47.1(c), first sentence)

To:
BRASNETT, Adrian, H.
Mewburn Ellis
York House
23 Kingsway
London WC2B 6HP
ROYAUME-UNI

RECEIVED
21 JAN 2000

Date of ~~mailing~~ (day/month/year) 13 January 2000 (13.01.00)	
Applicant's or agent's file reference AHB/CP5775333	**IMPORTANT NOTICE**

International application No.	**International filing date** (day/month/year)	**Priority date** (day/month/year)
~~PCT~~/GB99/02123	02 July 1999 (02.07.99)	02 July 1998 (02.07.98)

Applicant
CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED et al

1. Notice is hereby given that the International Bureau has communicated, as provided in Article 20, the international application to the following designated Offices on the date indicated above as the date of mailing of this Notice:
   AU,CN,EP,IL,JP,KP,KR,US

   In accordance with Rule 47.1(c), third sentence, those Offices will accept the present Notice as conclusive evidence that the communication of the international application has duly taken place on the date of mailing indicated above and no copy of the international application is required to be furnished by the applicant to the designated Office(s).

2. The following designated Offices have waived the requirement for such a communication at this time:
   AE,AL,AM,AP,AT,AZ,BA,BB,BG,BR,BY,CA,CH,CU,CZ,DE,DK,EA,EE,ES,FI,GB,GD,GE,GH,GM,HR,
   HU,ID,IN,IS,KE,KG,KZ,LC,LK,LR,LS,LT,LU,LV,MD,MG,MK,MN,MW,MX,NO,NZ,OA,PL,PT,RO,RU,
   SD,SE,SG,SI,SK,SL,TJ,TM,TR,TT,UA,UG,UZ,VN,YU,ZA,ZW
   The communication will be made to those Offices only upon their request. Furthermore, those Offices do not require the applicant to furnish a copy of the international application (Rule 49.1(a-bis)).

3. Enclosed with this Notice is a copy of the international application as published by the International Bureau on
   13 January 2000 (13.01.00) under No. WO 00/01822

REMINDER REGARDING CHAPTER II (Article 31(2)(a) and Rule 54.2)

If the applicant wishes to postpone entry into the national phase until 30 months (or later in some Offices) from the priority date, **a demand for international preliminary examination** must be filed with the competent International Preliminary Examining Authority before the expiration of 19 months from the priority date.

It is the applicant's sole responsibility to monitor the 19-month time limit.

Note that only an applicant who is a national or resident of a PCT Contracting State which is bound by Chapter II has the right to file a demand for international preliminary examination.

REMINDER REGARDING ENTRY INTO THE NATIONAL PHASE (Article 22 or 39(1))

If the applicant wishes to proceed with the international application in the **national phase**, he must, within 20 months or 30 months, or later in some Offices, perform the acts referred to therein before each designated or elected Office.

For further important information on the time limits and acts to be performed for entering the national phase, see the Annex to Form PCT/IB/301 (Notification of Receipt of Record Copy) and Volume II of the PCT Applicant's Guide.

The International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland

Facsimile No. (41-22) 740.14.35

Authorized officer
J. Zahra

Telephone No. (41-22) 338.83.38

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

PCT

To:

BRASNETT, Adrian H.
Mewburn Ellis
York House
23 Kingsway
London WC2B 6HP
GRANDE BRETAGNE

RECEIVED
17 FEB 2000

NOTIFICATION OF RECEIPT OF DEMAND BY COMPETENT INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

(PCT Rules 59.3(e) and 61.1(b), first sentence and Administrative Instructions, Section 601(a))

Date of mailing *(day/month/year)*	15.02.00

Applicant's or agent's file reference	IMPORTANT NOTIFICATION
AHB/CP5775333	

International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB 99/02123	02/07/1999	02/07/1998

Applicant

CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED et al.

1. The applicant is hereby **notified** that this International Preliminary Examining Authority considers the following date as the date of receipt of the demand for international preliminary examination of the international application:

01/02/2000

2. This date of receipt is:

- [x] the actual date of receipt of the demand by this Authority (Rule 61.1(b)).
- [ ] the actual date of receipt of the demand on behalf of this Authority (Rule 59.3(e)).
- [ ] the date on which this Authority has, in response to the invitation to correct defects in the demand (Form PCT/IPEA/404), received the required corrections.

3. [ ] **ATTENTION:** That date of receipt is **AFTER** the expiration of 19 months from the priority date. Consequently, the election(s) made in the demand does (do) not have the effect of postponing the entry into the national phase until 30 months from the priority date (or later in some Offices) (Article 39(1)). Therefore, the acts for entry into the national phase must be performed within 20 months from the priority date (or later in some Offices) (Article 22). For details, see the *PCT Applicant's Guide*, Volume II.

[ ] *(If applicable)* This notification confirms the information given by telephone, facsimile transmission or in person on: _____

4. Only where paragraph 3 applies, a copy of this notification has been sent to the International Bureau.

Name and mailing address of the IPEA/	Authorized officer
European Patent Office D-80298 Munich Tel. (+49-89) 2399-0, Tx: 523656 epmu d Fax: (+49-89) 2399-4465	CHAVONAND F H Tel. (+49-89) 2399-2390

EUROPÄISCHES PATENTAMT · EUROPEAN PATENT OFFICE · OFFICE EUROPÉEN DES BREVETS

PATENT COOPERATION TREATY

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

BRASNETT, Adrian H.
Mewburn Ellis
York House
23 Kingsway
London WC2B 6HP
GRANDE BRETAGNE

PCT

NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL PRELIMINARY EXAMINATION REPORT
(PCT Rule 71.1)

Date of mailing *(day/month/year)*	05.10.2000
Applicant's or agent's file reference: AHB/CP5775333	**IMPORTANT NOTIFICATION**

International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB99/02123	02/07/1999	02/07/1998

Applicant
CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED et al.

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.

2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.

3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. **REMINDER**

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/	Authorized officer
European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d	Vullo, C

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
AHB/CP5775333		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB99/02123	02/07/1999	02/07/1998
International Patent Classification (IPC) or national classification and IPC C12N15/13		
Applicant CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 8 sheets, including this cover sheet.

   ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☒ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☐ Certain documents cited
   - VII ☐ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
01/02/2000	05.10.2000
Name and mailing address of the international preliminary examining authority: European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d	Authorized officer Vix, O

IN RNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/02123

I. Basis of the r port

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

Description, pages:

1-41 as originally filed

Claims, No.:

1-16 as originally filed

Drawings, sheets:

1/8-8/8 as originally filed

2. The amendments have resulted in the cancellation of:

☐ the description, pages:
☐ the claims, Nos.:
☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

IV. Lack of unity of invention

1. In response to the invitation to restrict or pay additional fees the applicant has:

☐ restricted the claims.

☐ paid additional fees.

☐ paid additional fees under protest.

☐ neither restricted nor paid additional fees.

2. ☒ This Authority found that the requirement of unity of invention is not complied and chose, according to Rule 68.1, not to invite the applicant to restrict or pay additional fees.

3. This Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3 is

   ☐ complied with.

   ☒ not complied with for the following reasons:

   see separate sheet

4. Consequently, the following parts of the international application were the subject of international preliminary examination in establishing this report:

   ☒ all parts.

   ☐ the parts relating to claims Nos. .

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	9-16
	No:	Claims	1-8
Inventive step (IS)	Yes:	Claims	
	No:	Claims	1-16
Industrial applicability (IA)	Yes:	Claims	1-15
	No:	Claims	16 (see item V)

2. Citations and explanations

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate she t

Reference is made to the following documents:

D1: MILLER G K ET AL: 'Immunologic and biochemical analysis of TNT-1 and TNT-2 monoclonal antibody binding to histones.' HYBRIDOMA, (1993 DEC) 12 (6) 689-98. , XP002119491 cited in the application

D2: KING D J ET AL: 'PREPARATION AND PRECLINICAL EVALUATION OF HUMANISED A33 IMMUNOCONJUGATES FOR RADIOIMMUNOTHERAPY' BRITISH JOURNAL OF CANCER, vol. 72, no. 6, December 1995 (1995-12), pages 1364-1372, XP002911796 ISSN: 0007-0920

D3: DATABASE EMBL, PIR2: 'ACCESSION NUMBER: S47184' RESIDUES 1-109,6 June 1995 (1995-06-06), XP002119493

D4: DATABASE EMBL, PIR2: 'Accession number: S19663' RESIDUES 1-109, XP002119494

D5: DESRUES B ET AL: 'Monoclonal antibody Po66 uptake by human lung tumours implanted in nude mice: effect of co-administration with doxorubicin.' BRITISH JOURNAL OF CANCER, (1995 NOV) 72 (5) 1076-82. , XP002119492

D6: US-A-5 019 368 (EPSTEIN ALAN L ET AL) 28 May 1991 (1991-05-28)

Re Item IV

Lack of unity of invention

The common inventive concept between all claims is the intracellular antigen binding capacity of a "binding member" having an antibody framework or part thereof. Such concept is not new or, if at all new by the presence of specific Complementary Determining Regions sequences -CDRs-, it is not inventive in view of D1. D1 refers to monoclonal antibodies (mAb) binding to intracellular antigens (such as H1 and H3 histones) and their use as reagents for the diagnosis and treatment of cancer.

Therefore the binding members of independent claim 1 and 4 are not anymore linked by said common inventive concept as to form a single valid invention (as required by Rule 13.1 PCT).

Correspondingly, present claims do not relate to one invention but to two separate ones, namely:

- invention 1 : the antibody or part thereof comprising a polypeptide domain (or CDR3) comprising residues 99 to 106 of SEQ ID N°2 (claims 1-4 complete, claims 9-16 partial).
- invention 2 : the antibody or part thereof comprising a polypeptide domain comprising residues 88 to 98 of SEQ ID N°4 (claims 5-8 complete, claims 9-16 partial).

Re Item V

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

The application relates to antibodies or fragment thereof which bind to necrotic centre of tumors (binding to nuclear proteins like Histones), and their use in treatment of cancer and diagnosis (radioactive or toxic labelled antibody).

1. Novelty (Art. 33(2) PCT)

1.1 Both D3 and D4 disclose a human variable chain domain sequence showing a 100% identity with the sequence ID N°4 of the present application. Thus D3 and D4 are prejudicial for novelty of claims 5-8 dealing with a polypeptide binding domain comprising parts of sequence ID N°4.
Consequently, claims 5-8 do not satisfy the criterion set forth in Article 33(2) PCT.

1.2 From D6 it is known that several antibodies directed to an internal cellular component have been obtained and used for diagnosis and treatment of tumours.
Claims 1-4 refer to a broad and vague definition of the polypeptide binding domain. This binding molecule is defined as "comprising" a sequence "substantially" as set out in the CDRs regions of SEQ ID N°2. It cannot be excluded that the sequence of the monoclonal antibodies binding to intracellular antigen presented in Table 1 (column 10-11) of D6 fall under the scope of claims 1-4 in particular in view of the term "substantially comprising" allowing variations of the determinants of SEQ ID N°2. Therefore, claims 1-4 lack novelty in view of D6.

2. Inventive step (Art. 33(3) PCT)

The present application deals with "binding elements" derived from an antibody framework (like scFv or $F(ab')_2$) that are binding to intracellular antigen. These binding elements are used for treatment and diagnosis of tumour in the human body.

Rendering the above objected claims novel, the technical problem to be solved by the present invention (in view of the available prior art, e.g. D1/D6) may be regarded as the provision of further antibodies, or fragments thereof, binding to intracellular antigens and useful for the diagnosis and treatment of cancer.

The problem was solved by the present application using different repertoire for CDR variation within the antibody antigen binding site.

Such solution is based on a known technique in the art of immunology and immunochemistry : as the antibody recognition is based on the CDRs regions, a standard strategy for the man skilled in the art is to screen antibody repertoires for selection of new antibodies showing a specificity for the antigen of interest.
The use of labelled antibodies for tumour imaging or treatment is also well known in the art as seen in D5/D6 (in addition to D1) which disclose examples of labelled antibodies production (radiolabels, conjugation with toxic or pharmaceutical compounds, etc...- see D6 column 3-6 and D5 page 1081).
Moreover, concerning the dependent claim 3 and the concept of humanisation, according to D2, the use of radioimmunoconjugates derived from rodents for tumour treatment in patients, and the related problem of developing a human anti-mouse antibody response (HAMA) were known to the person skilled in the art (see D2 page 1364). A solution provided by D2 is the preparation of new humanised radioimmunoconjugates for treatment of tumours. The technique of antibody "humanisation" is well known for the person of skill : a human antibody or part of it can be used as a framework to graft the murine CDRs parts responsible for the specific tumour antigen binding. The clinical studies reported in D2 (page 1371) also suggest that the humanised antibodies have a longer circulating half-life compared to the murine ones, detail which becomes important when using radioimmunotherapy methods.

The variation of size of a "binding member" derived from an antibody is also widely used and based on routine methods in the field of immunology : the structural features important for antigen recognition are formed by the interaction of the light and heavy variable domains (allowing the 3 CDRs to be spatially close and form the antigen binding pocket). Therefore, any molecule that possess the two variable domains VH and VL, each carrying the 3 appropriate CDRs, will conserve the binding capacities of the intact antibody. Consequently, the provision of antibody derived structural elements like the known Fab, F(ab')2, Fv or single chain Fv (scFv) are not considered to involve an inventive step.

In consequence, in view of the documents D1 -or also D5/D6- (and D2 for the humanisation concept) the man skilled in the art would arrive to the claimed subject-matter of the invention without applying inventive activity. Thus , claims 1-16 do not satisfy the criterion set forth in Article 33(3) PCT.

3. Industrial applicability (Art. 33(4) PCT)

For the assessment of the present claims 16 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. Consequently, under the provision of Rule 67.1(iv) PCT, no statement with regard to industrial applicability of said claims will be made (Article 34(4)(a)(i) PCT).

Re Item VIII

Certain observations on the international application

1. The term "isolated specific binding member", present in the majority of the claims, is vague and open to interpretation when lacking precise technical features (size, sequence, etc...). It is not clear whether it refers strictly to a polypeptide or any other type of hybrid molecule simply "comprising a polypeptide binding domain". From the description and the type of sequence listed, these functional domains are all related to an antibody framework carrying three hypervariable loops or Complementary Determining Regions -CDRs- responsible of the binding (either partial antibody

functional element like $F(ab')_2$ or Fab, etc...). This support in the description does not include any possible hypothetical binding member and should be restricted to an antibody-antigen system (as discussed in page 5).

2. The wording "substantially" in claims 1-2, 4-6, 8 and 15 refers to a relative definition open to interpretation, and thus render the scope of said claims unclear (Article 6 PCT).

3. Claims 1-4 lack additional meaningful technical characterising features. The binding capacity of these short sequences are only possible if they have the appropriate positions in the hypervariable loops on an antibody framework. Therefore, the antibody framework is an important characterising feature of the invention and should be included in the independant claims.

4. The expression "intracellular antigen" in claims 1 and 5 is vague and open to interpretation. Due to the very high cellular protein content considered as potential antigen, it is not clear what type of antigen might be considered to fall within the scope of said claims.

5. F(ab')2 and scFv should be clearly defined in claim 10.

6. Claim 11 relates to "detectable and functional label" without any characterising features. Such a claim is open to interpretation and appears unclear since there is no mention what type of detection technique or functional properties might be involved in said claim.

7. The tumour:blood ratio mentioned in claim 15d) might depend on the calculation method being used. Such ratio might therefore be ambiguous and lack clarity.

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
AHB/CP5775333		
International application No.	International filing date *(day/month/year)*	(Earliest) Priority Date *(day/month/year)*
PCT/GB 99/ 02123	02/07/1999	02/07/1998
Applicant		
CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED et al.		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 6 sheets.

[X] It is also accompanied by a copy of each prior art document cited in this report.

1. **Basis of the report**

a. With regard to the **language,** the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

[ ] the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

[X] contained in the international application in written form.

[X] filed together with the international application in computer readable form.

[ ] furnished subsequently to this Authority in written form.

[ ] furnished subsequently to this Authority in computer readble form.

[ ] the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

[ ] the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. [X] **Certain claims were found unsearchable** (See Box I).

3. [ ] **Unity of invention is lacking** (see Box II).

4. With regard to the **title,**

[X] the text is approved as submitted by the applicant.

[ ] the text has been established by this Authority to read as follows:

5. With regard to the **abstract,**

[X] the text is approved as submitted by the applicant.

[ ] the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. _____

[ ] as suggested by the applicant.

[ ] because the applicant failed to suggest a figure.

[ ] because this figure better characterizes the invention.

[X] None of the figures.

INTERNATIONAL SEARCH REPORT

ernational application No.
PCT/GB 99/02123

Box I Observations where certain claims wer found unsearchable (Continuation of item 1 of first she t)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. [X] Claims Nos.: 16
because they relate to subject matter not required to be searched by this Authority, namely:

see FURTHER INFORMATION sheet PCT/ISA/210

2. [ ] Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. [ ] Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. [ ] As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. [ ] As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. [ ] No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 16 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/ composition.

Continuation of Box I.1

Claims Nos.: 16

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 99/02123

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C12N15/13 C12N15/62 C07K16/18 C07K16/46 G01N33/577
A61K51/10 A61K39/395 //G01N33/574

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07K G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	MILLER G K ET AL: "Immunologic and biochemical analysis of TNT-1 and TNT-2 monoclonal antibody binding to histones." HYBRIDOMA, (1993 DEC) 12 (6) 689-98. , XP002119491 cited in the application abstract --- -/--	1-16

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
20 October 1999	04/11/1999

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016	Covone, M

Form PCT/ISA/210 (second sheet) (July 1992)

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication,where appropriate, of the relevant passages	Relevant to claim No.
Y	KING D J ET AL: "PREPARATION AND PRECLINICAL EVALUATION OF HUMANISED A33 IMMUNOCONJUGATES FOR RADIOIMMUNOTHERAPY" BRITISH JOURNAL OF CANCER, vol. 72, no. 6, December 1995 (1995-12), pages 1364-1372, XP002911796 ISSN: 0007-0920 page 1364, right-hand column, paragraph 3 -page 1365, left-hand column, paragraph 2 page 1369, right-hand column, paragraphs 2,3 page 1371, right-hand column, paragraph 3 ---	1-16
X	DATABASE BIOSIS 'Online! BIOSCIENCES INFORMATION SERVICE, PHILADELPHIA, PA, US AKIYAMA, YOSHIMITSU (1) ET AL: "Cell Proliferation Kinetics of Human Gastric Carcinoma -A Quantitative Study of PCNA Expression Using a Color Image Analysis System." retrieved from STN XP002119497 abstract & STOMACH AND INTESTINE, (1995) VOL. 30, NO. 1, PP. 113-119. , ---	1,2,4-6, 8,9,11, 14,16
X	DESRUES B ET AL: "Monoclonal antibody Po66 uptake by human lung tumours implanted in nude mice: effect of co-administration with doxorubicin." BRITISH JOURNAL OF CANCER, (1995 NOV) 72 (5) 1076-82. , XP002119492 abstract page 1076, right-hand column, paragraph 2 page 1077, left-hand column, line 1 page 1080, right-hand column, paragraph 1 page 1081, right-hand column, paragraph 1 ---	1,2,4-6, 8-11,14, 16
X	US 5 019 368 A (EPSTEIN ALAN L ET AL) 28 May 1991 (1991-05-28) column 2, line 62 -column 3, line 2 column 4, line 31-42 column 5, line 25-52 column 7, line 25-52 claims ---	1,2,4-6, 8,9,11, 14,16
X	DATABASE EMBL, PIR2: "ACCESSION NUMBER: S47184" RESIDUES 1-109,6 June 1995 (1995-06-06), XP002119493 the whole document ---	5-8

-/--

1

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication,where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE EMBL, PIR2: "Accession number: S19663" RESIDUES 1-109, XP002119494 the whole document ---	5-8
A	DATABASE EMBL, EMHUM3: "ACCESSION NUMBER: Z14195, entry Hse54427 " RESIDUES 1-434, 20 January 1993 (1993-01-20), XP002119495 the whole document ---	15
P,X	HORNICK J.L. ET AL: "A new chemically modified chimeric TNT-3 monoclonal antibody directed against DNA for the radioimmunotherapy of solid tumors." CANCER BIOTHERAPY AND RADIOPHARMACEUTICALS, (1998) 13/4 (255-268). , XP002119496 abstract page 256, left-hand column, paragraph 2 -right-hand column, paragraph 1 page 257, right-hand column, paragraph 3 -page 258, left-hand column, paragraph 3 page 266, left-hand column, paragraph 3 -page 267, left-hand column, paragraph 1 -----	1-16

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 99/02123

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5019368 A	28-05-1991	US 5882626 A	16-03-1999

M.H

PCT WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 :
C12N 15/13, 15/62, C07K 16/18, 16/46, G01N 33/577, A61K 51/10, 39/395 // G01N 33/574

A1

(11) International Publication Number: **WO 00/01822**

(43) International Publication Date: 13 January 2000 (13.01.00)

(21) International Application Number: PCT/GB99/02123

(22) International Filing Date: 2 July 1999 (02.07.99)

(30) Priority Data:
9814383.7 2 July 1998 (02.07.98) GB

(71) Applicant *(for all designated States except US)*: CAMBRIDGE ANTIBODY TECHNOLOGY LIMITED [GB/GB]; The Science Park, Melbourn, Royston, Cambridgeshire SG8 6JJ (GB).

(72) Inventors; and
(75) Inventors/Applicants *(for US only)*: WILLIAMS, Andrew, James [GB/GB]; Larksmead, Long Lane, Fowlmere, Royston SG8 7TG (GB). TEMPEST, Philip, Ronald [GB/GB]; 43 High Street, West Wratting, Cambridgeshire CB1 5LU (GB). HOLTET, Thor, Las [DK/DK]; Morkhoj Bygade 16, DK–2860 Soborg (DK). JACKSON, Helen [GB/GB]; Peace Cottage, 15 Bedford Road, North Hill, Bedfordshire SG18 9AH (GB).

(74) Agents: BRASNETT, Adrian, H. et al.; Mewburn Ellis, York House, 23 Kingsway, London WC2B 6HP (GB).

(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published
With international search report.
Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: SPECIFIC BINDING PROTEINS INCLUDING ANTIBODIES WHICH BIND TO THE NECROTIC CENTRE OF TUMOURS, AND USES THEREOF

(57) Abstract

Specific binding members, based on the third CDR of the antibody NHS76 (SEQ ID NO: 2) are provided, together with their use in methods of treatment and diagnosis.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

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SPECIFIC BINDING PROTEINS INCLUDING ANTIBODIES WHICH BIND TO THE NECROTIC CENTRE OF TUMOURS, AND USES THEREOF.

The present invention relates to specific binding members, particularly antibodies and fragments thereof, which bind to the necrotic centres of tumours. These specific binding members are useful in the treatment of cancer.

Background to the Invention.

The treatment of proliferative disease, particularly cancer, by chemotherapeutic means often relies upon exploiting differences in target proliferating cells and other normal cells in the human or animal body. For example, many chemical agents are designed to be taken up by rapidly replicating DNA so that the process of DNA replication and cell division is disrupted. Another approach is to identify antigens on the surface of tumour cells which are not normally expressed in developed human tissue, such as embryonic antigens. Such antigens can be targeted with binding proteins such as antibodies to deliver a toxic agent to or other substance which is capable directly or indirectly of activating a toxic agent at, the site of a tumour.

An alternative approach which has been developed more recently relies on the fact that a significant proportion of tumour cells are in various stages of cell degeneration and death. Unlike programmed cell death (apoptosis) which occurs during the natural turnover of certain cell types, tumour cells undergo a degenerative and less controlled death during which they have been found to exhibit abnormal surface membrane permeability. European patent application EP-A-270340 describes the preparation of murine antibodies to nuclear components of cells which are able to target necrotic cells in a tumour by exploiting this phenomenon. Miller et al (Hybridoma, 1993, 12, 689-697) describe a particular antibody prepared in accordance with EP-A-270340. This antibody, TNT-1,

was found to bind histone fraction H1. The histones are the most abundant proteins in chromatin, the concentration of each type of histone being approximately 5000 times higher than the concentration of a typical sequence-specific DNA-binding protein. Epstein et al (in "Handbook of Targeted Delivery of Imaging Agents" CRC press, Inc., 1995, ed. VP Torchilin) describes the uptake of radiolabelled TNT-1 into nude mice bearing the ME-180 cervical carcinoma, and report that there was no appreciable labelling of other organs. The authors also used the $F(ab')_2$ fragment of TNT-1 labelled with ^{131}I for tumour imaging studies in human patients.

A known problem with the use of murine antibodies in human therapy is that repeat treatment of such antibodies leads to a human anti-mouse antibody (HAMA) response. HAMA responses have a range of effects, from neutralisation of the administered antibody leading to a reduced therapeutic dose, through to allergic responses, serum sickness and renal impairment. In order to overcome these disadvantages humanization of antibodies has been developed. More recently, repertoires of human antibodies have been cloned and these can be screened, for example by phage display technology, (McCafferty et al, WO92/01047) to identify human antibodies specific for human antigens.

Disclosure of the Invention

We have investigated the binding properties of the TNT-1 antibody and attempted to secure human antibodies with similar binding profiles and useful tumour localisation in animal models. We have found that although TNT-1 binds primarily to nuclear histone H1 it also binds a complex pattern of other histones and non-identified proteins in a nuclear preparation. We were able to identify a number of human antibodies, screened primarily against a nuclear extract and further against histone

H1 which had similar binding profiles. Despite the similarities in binding profiles between the human antibody clones analysed, only one (identified below as "NHS76") was found to additionally exhibit low cross reactivity to non-necrotic tissues and organs and good tumour:blood localisation ratio. Other candidate antibodies tested either showed cross-reactivity or had poorer tumour:blood ratios of localisation.

Accordingly, specific binding proteins such as antibodies which are based on the complementarity-determining regions (CDRs) of the NHS76 antibody identified, particularly the CDR 3 regions, will be useful for targeting the necrotic centres of malignant tumours.

In the accompanying drawings, the nucleic acid sequence and translation thereof of the NHS76 VH gene is shown in Figure 1. The nucleic acid sequence is designated SEQ ID NO:1. The translation is SEQ ID NO:2. The VL gene of NHS 76 is shown as Figure 2. Its nucleic acid sequence is designated SEQ ID NO:3, and its translation as SEQ ID NO:4. In both figures, the CDR's are indicated in boxes.

In a first aspect the present invention provides an isolated specific binding member capable of binding an intracellular antigen, wherein said specific binding member comprises a polypeptide binding domain comprising an amino acid sequence substantially as set out as residues 99 to 106 of SEQ ID NO:2. The invention further provides said isolated specific binding member which further comprises one or both of the polypeptide binding domains substantially as set out as residues 31-36 and 51-66 of SEQ ID NO:2, preferably both. In a preferred embodiment, the binding domains are carried by a human antibody framework. One example of such an embodiment is the sequence substantially as shown in SEQ ID NO:2.

In a second aspect, the invention provides an isolated specific binding member capable of binding an intracellular antigen, wherein said specific binding member comprises a polypeptide binding domain comprising an amino acid sequence substantially as set out as residues 88 to 98 of SEQ ID NO:4. The invention further provides said isolated specific binding member which further comprises one or both of the polypeptide binding domains substantially as set out as residues 23-33 and 49-55 of SEQ ID NO:4, preferably both. In a preferred embodiment, the binding domains are carried by a human antibody framework. One example of such an embodiment is the sequence substantially as shown in SEQ ID NO:4.

In a particularly preferred embodiment, the invention provides a specific binding member which comprises a first specific binding member comprising a sequence substantially as set out as residues 99 to 106 of SEQ ID NO:2 in association with a second specific binding member comprising a sequence substantially set out as residues 88 to 98 of SEQ ID NO:4. Such a specific binding member according to the invention may be in the form of an antibody F(ab')2 fragment.

Specific binding members of the invention may carry a detectable or functional label.

In further aspects, the invention provides an isolated nucleic acid which comprises a sequence encoding a specific binding member as defined above, and methods of preparing specific binding members of the invention which comprise expressing said nucleic acids under conditions to bring about expression of said binding member, and recovering the binding member.

Specific binding members according to the invention may be used in a method of treatment or diagnosis of the human or animal body, such as a method of treatment of a tumour in a human

patient which comprises administering to said patient an effective amount of a specific binding member of the invention.

These and other aspects of the invention are described in further detail below.

Detailed Description of the Invention.

A. TERMINOLOGY

Specific binding member

This describes a member of a pair of molecules which have binding specificity for one another. The members of a specific binding pair may be naturally derived or wholly or partially synthetically produced. One member of the pair of molecules has an area on its surface, or a cavity, which specifically binds to and is therefore complementary to a particular spatial and polar organisation of the other member of the pair of molecules. Thus the members of the pair have the property of binding specifically to each other. Examples of types of specific binding pairs are antigen-antibody, biotin-avidin, hormone-hormone receptor, receptor-ligand, enzyme-substrate. This application is concerned with antigen-antibody type reactions.

Antibody

This describes an immunoglobulin whether natural or partly or wholly synthetically produced. The term also covers any polypeptide or protein having a binding domain which is, or is homologous to, an antibody binding domain. These can be derived from natural sources, or they may be partly or wholly synthetically produced. Examples of antibodies are the immunoglobulin isotypes and their isotypic subclasses; fragments which comprise an antigen binding domain such as Fab, scFv, Fv, dAb, Fd; and diabodies.

It is possible to take monoclonal and other antibodies and use techniques of recombinant DNA technology to produce other antibodies or chimeric molecules which retain the specificity of the original antibody. Such techniques may involve introducing DNA encoding the immunoglobulin variable region, or the complementarity determining regions (CDRs), of an antibody to the constant regions, or constant regions plus framework regions, of a different immunoglobulin. See, for instance, EP-A-184187, GB 2188638A or EP-A-239400. A hybridoma or other cell producing an antibody may be subject to genetic mutation or other changes, which may or may not alter the binding specificity of antibodies produced.

As antibodies can be modified in a number of ways, the term "antibody" should be construed as covering any specific binding member or substance having a binding domain with the required specificity. Thus, this term covers antibody fragments, derivatives, functional equivalents and homologues of antibodies, including any polypeptide comprising an immunoglobulin binding domain, whether natural or wholly or partially synthetic. Chimeric molecules comprising an immunoglobulin binding domain, or equivalent, fused to another polypeptide are therefore included. Cloning and expression of chimeric antibodies are described in EP-A-0120694 and EP-A-0125023.

It has been shown that fragments of a whole antibody can perform the function of binding antigens. Examples of binding fragments are (i) the Fab fragment consisting of VL, VH, CL and CH1 domains; (ii) the Fd fragment consisting of the VH and CH1 domains; (iii) the Fv fragment consisting of the VL and VH domains of a single antibody; (iv) the dAb fragment (Ward, E.S. et al., Nature 341, 544-546 (1989)) which consists of a VH domain; (v) isolated CDR regions; (vi) F(ab')2 fragments, a bivalent fragment comprising two linked Fab fragments (vii)

single chain Fv molecules (scFv), wherein a VH domain and a VL domain are linked by a peptide linker which allows the two domains to associate to form an antigen binding site (Bird et al, Science, **242**, 423-426, 1988; Huston et al, PNAS USA, **85**, 5879-5883, 1988); (viii) bispecific single chain Fv dimers (PCT/US92/09965) and (ix) "diabodies", multivalent or multispecific fragments constructed by gene fusion (WO94/13804; P. Holliger et al Proc. Natl. Acad. Sci. USA **90** 6444-6448, 1993).

Diabodies are multimers of polypeptides, each polypeptide comprising a first domain comprising a binding region of an immunoglobulin light chain and a second domain comprising a binding region of an immunoglobulin heavy chain, the two domains being linked (e.g. by a peptide linker) but unable to associate with each other to form an antigen binding site: antigen binding sites are formed by the association of the first domain of one polypeptide within the multimer with the second domain of another polypeptide within the multimer (WO94/13804).

Where bispecific antibodies are to be used, these may be conventional bispecific antibodies, which can be manufactured in a variety of ways (Holliger, P. and Winter G. Current Opinion Biotechnol. **4**, 446-449 (1993)), eg prepared chemically or from hybrid hybridomas, or may be any of the bispecific antibody fragments mentioned above. It may be preferable to use scFv dimers or diabodies rather than whole antibodies. Diabodies and scFv can be constructed without an Fc region, using only variable domains, potentially reducing the effects of anti-idiotypic reaction. Other forms of bispecific antibodies include the single chain "Janusins" described in Traunecker et al, EMBO Journal, 10, 3655-3659, (1991).

Bispecific diabodies, as opposed to bispecific whole

antibodies, may also be particularly useful because they can be readily constructed and expressed in *E.coli*. Diabodies (and many other polypeptides such as antibody fragments) of appropriate binding specificities can be readily selected using phage display (WO94/13804) from libraries. If one arm of the diabody is to be kept constant, for instance, with a specificity directed against antigen X, then a library can be made where the other arm is varied and an antibody of appropriate specificity selected.

Antigen binding domain

This describes the part of an antibody which comprises the area which specifically binds to and is complementary to part or all of an antigen. Where an antigen is large, an antibody may only bind to a particular part of the antigen, which part is termed an epitope. An antigen binding domain may be provided by one or more antibody variable domains. Preferably, an antigen binding domain comprises an antibody light chain variable region (VL) and an antibody heavy chain variable region (VH).

Specific

This may be used to refer to the situation in which one member of a specific binding pair will not show any significant binding to molecules other than its specific binding partner(s). The term is also applicable where e.g. an antigen binding domain is specific for a particular epitope which is carried by a number of antigens, in which case the specific binding member carrying the antigen binding domain will be able to bind to the various antigens carrying the epitope.

Comprise

This is generally used in the sense of include, that is to say permitting the presence of one or more features or components.

Isolated

This refers to the state in which specific binding members of the invention, or nucleic acid encoding such binding members will be, in accordance with the present invention. Members and nucleic acid will be free or substantially free of material with which they are naturally associated such as other polypeptides or nucleic acids with which they are found in their natural environment, or the environment in which they are prepared (e.g. cell culture) when such preparation is by recombinant DNA technology practised *in vitro* or *in vivo*. Members and nucleic acid may be formulated with diluents or adjuvants and still for practical purposes be isolated - for example the members will normally be mixed with gelatin or other carriers if used to coat microtitre plates for use in immunoassays, or will be mixed with pharmaceutically acceptable carriers or diluents when used in diagnosis or therapy. Specific binding members may be glycosylated, either naturally or by systems of heterologous eukaryotic cells, or they may be (for example if produced by expression in a prokaryotic cell) unglycosylated.

B. DETAILED DISCLOSURE.

Isolated specific binding members of the invention are those such as NHS76 which are capable of binding an intracellular antigen. In particular, the member will bind to human histone H1. However the NHS76 antibody which forms a particular aspect of the invention also shows a pattern of reactivity to a range of intracellular antigens. It is preferred that other binding members of the invention also show the same or substantially similar pattern reactivity. This may be determined by comparing such members with an antibody comprising the VH and VL domains shown in SEQ ID NO:2 and SEQ ID NO:4 respectively. The comparison will typically be made using a western blot in which binding members are bound to duplicate blots prepared from a nuclear preparation of cells so that the pattern of

binding can be directly compared. Suitable nuclear preparations and conditions for western blotting are described below in the accompanying examples.

In particular, NHS76 shows differential binding to histones. This is reflected in the ELISA data of histone binding (Example 7 below) which shows that NHS shows greatest binding to histone H1 followed by H3 > H2B > H2A, H4. Suitable western blot conditions for measurement of binding to histones are illustrated in the accompanying examples.

As outlined above, we have found that the NHS76 antibody has *in vivo* properties which are not shared by other antibodies with apparently similar binding profiles. While not wishing to be bound by any one particular theory, one explanation for this is that the specific antibody-antigen interaction of the antibody results in the unexpectedly desirable *in vivo* activity. The binding of an antibody to its target antigen is mediated through the CDRs of its heavy and light chains, with the role of CDR3 being of particular importance. Accordingly, specific binding members based on the CDR3 regions of the heavy or light chain, and preferably both, of NHS76 will be useful specific binding members for *in vivo* therapy.

In general, the CDR3 regions, comprising amino acid sequences substantially as set out as residues 99 to 106 of SEQ ID NO:2 and 88 to 98 of SEQ ID NO:4 will be carried in a structure which allows for binding of the CDR3 regions to an intracellular antigen.

By "substantially as set out" it is meant that the CDR3 regions of the invention will be either identical or highly homologous to the specified regions of SEQ ID NO:2 and SEQ ID NO:4. By "highly homologous" it is contemplated that from 1 to 5, preferably from 1 to 4 such as 1 to 3 or 1 or 2 substitutions

may be made in the CDRs.

The structure for carrying the CDR3s of the invention will generally be of an antibody heavy or light chain sequence or substantial portion thereof in which the CDR3 regions are located at locations corresponding to the CDR3 region of naturally occurring VH and VL antibody variable domains encoded by rearranged immunoglobulin genes. The structures and locations of immunoglobulin variable domains may be determined by reference to (Kabat, E.A. et al, Sequences of Proteins of Immunological Interest. 4th Edition. US Department of Health and Human Services. 1987, and updates thereof, now available on the Internet (http://immuno.bme.nwu.edu)).

Preferably, the amino acid sequence substantially as set out as residues 99 to 106 of SEQ ID NO:2 is carried as the CDR3 in a human heavy chain variable domain or a substantial portion thereof, and the amino acid sequence substantially as set out as residues 88 to 98 of SEQ ID NO:4 is carried as the CDR3 in a human light chain variable domain or a substantial portion thereof.

The variable domains may be derived from any germline or rearranged human variable domain, or may be a synthetic variable domain based on consensus sequences of known human variable domains. The CDR3-derived sequences of the invention, as defined in the preceding paragraph, may be introduced into a repertoire of variable domains lacking CDR3 regions, using recombinant DNA technology.

For example, Marks *et al* (*Bio/Technology*, 1992, 10:779-783) describe methods of producing repertoires of antibody variable domains in which consensus primers directed at or adjacent to the 5' end of the variable domain area are used in conjunction with consensus primers to the third framework region of human

VH genes to provide a repertoire of VH variable domains lacking a CDR3. Marks *et al* further describe how this repertoire may be combined with a CDR3 of a particular antibody. Using analogous techniques, the CDR3-derived sequences of the present invention may be shuffled with repertoires of VH or VL domains lacking a CDR3, and the shuffled complete VH or VL domains combined with a cognate VL or VH domain to provide specific binding members of the invention. The repertoire may then be displayed in a suitable host system such as the phage display system of WO92/01047 so that suitable specific binding members may be selected. A repertoire may consist of from anything from 10^4 individual members upwards, for example from 10^6 to 10^8 or 10^{10} members.

Analogous shuffling or combinatorial techniques are also disclosed by Stemmer (*Nature*, 1994, 370:389-391), who describes the technique in relation to a β-lactamase gene but observes that the approach may be used for the generation of antibodies.

A further alternative is to generate novel VH or VL regions carrying the CDR3-derived sequences of the invention using random mutagenesis of, for example, the NHS76 VH or VL genes to generate mutations within the entire variable domain. Such a technique is described by Gram *et al* (1992, *Proc. Natl. Acad. Sci., USA*, 89:3576-3580), who used error-prone PCR.

Another method which may be used is to direct mutagenesis to CDR regions of VH or VL genes. Such techniques are disclosed by Barbas *et al*, (1994, *Proc. Natl. Acad. Sci., USA*, 91:3809-3813) and Schier *et al* (1996, *J. Mol. Biol.* 263:551-567).

All the above described techniques are known as such in the art and in themselves do not form part of the present invention. The skilled person will be able to use such techniques to provide specific binding members of the invention using routine

methodology in the art.

A substantial portion of an immunoglobulin variable domain will comprise at least the three CDR regions, together with their intervening framework regions. Preferably, the portion will also include at least about 50% of either or both of the first and fourth framework regions, the 50% being the C-terminal 50% of the first framework region and the N-terminal 50% of the fourth framework region. Additional residues at the N-terminal or C-terminal end of the substantial part of the variable domain may be those not normally associated with naturally occurring variable domain regions. For example, construction of specific binding members of the present invention made by recombinant DNA techniques may result in the introduction of - or C-terminal residues encoded by linkers introduced to facilitate cloning or other manipulation steps. Other manipulation steps include the introduction of linkers to join variable domains of the invention to further protein sequences including immunoglobulin heavy chains, other variable domains (for example in the production of diabodies) or protein labels as discussed in more detail below.

Although in a preferred aspect of the invention specific binding members comprising a pair of binding domains based on sequences substantially set out in SEQ ID NO:2 and SEQ ID NO:4 are preferred, single binding domains based on either of these sequences form further aspects of the invention. In the case of the binding domains based on the sequence substantially set out in SEQ ID NO:2, such binding domains may be used as targeting agents for intracellular antigens since it is known that immunoglobulin VH domains are capable of binding target antigens in a specific manner.

In the case of either of the single chain specific binding domains, these domains may be used to screen for complementary domains capable of forming a two-domain specific binding member

which has *in vivo* properties as good as or equal to the NHS 76 antibody disclosed herein.

This may be achieved by phage display screening methods using the so-called hierarchical dual combinatorial approach as disclosed in WO 92/01047 in which an individual colony containing either an H or L chain clone is used to infect a complete library of clones encoding the other chain (L or H) and the resulting two-chain specific binding member is selected in accordance with phage display techniques such as those described in that reference. This technique is also disclosed in Marks *et al, ibid.*

Specific binding members of the present invention may further comprise antibody constant regions or parts thereof. For example, specific binding members based on SEQ ID NO:4 may be attached at their C-terminal end to antibody light chain constant domains including human Cκ or Cλ chains, preferably Cλ chains. Similarly, specific binding members based on SEQ ID NO:2 may be attached at their C-terminal end to all or part of an immunoglobulin heavy chain derived from any antibody isotype, e.g. IgG, IgA, IgE and IgM and any of the isotype sub-classes, particularly IgG1 and IgG4. IgG1 is preferred.

The *in vivo* properties, particularly with regard to tumour:blood ratio and rate of clearance, of specific binding members of the invention will be comparable to NHS76. Following administration to a human or animal subject such a specific binding member will show a peak tumour to blood ration of > 3:1. Preferably at such a ratio the specific binding member will also have an organ to blood ratio of < 1:1 in organs away from the site of the tumour. These ratios exclude organs of catabolism and secretion of the administered specific binding member. Thus in the case of scFvs and Fabs (as shown in the accompanying examples), the binding members are secreted via

the kidneys and there is greater localisation here than other organs. In the case of whole IgGs, clearance will be at least in part, via the liver. The peak localisation ratio will normally be achieved between the 48 and 96 hours following administration of the specific binding member. More particularly, the ratio may be measured in a tumour xenograft of about 0.2 - 1.0 g formed subcutaneously in one flank of an athymic nude mouse.

Antibodies of the invention may be labelled with a detectable or functional label. Detectable labels include radiolabels such as ^{131}I or ^{99}Tc, which may be attached to antibodies of the invention using conventional chemistry known in the art of antibody imaging. Labels also include enzyme labels such as horseradish peroxidase. Labels further include chemical moieties such as biotin which may be detected via binding to a specific cognate detectable moiety, e.g. labelled avidin.

Functional labels include substances which are designed to be targeted to the site of a tumour to cause destruction of tumour tissue. Such functional labels include toxins such as ricin and enzymes such as bacterial carboxypeptidase or nitroreductase, which are capable of converting prodrugs into active drugs at the site of a tumour.

Antibodies of the present invention are designed to be used in methods of diagnosis and treatment of tumours in human or animal subjects, particularly solid tumours which have a necrotic centre. These tumours may be primary or secondary solid tumours of any type including, but not limited to, cervical, ovarian, prostate, lung, liver, pancreatic, colon and stomach tumours.

Antibodies of the present invention may be administered to a patient in need of treatment via any suitable route, usually by

injection into the bloodstream or directly into the site of the tumour. The precise dose will depend upon a number of factors, including whether the antibody is for diagnosis or for treatment, the size and location of the tumour, the precise nature of the antibody (e.g. whole antibody, fragment or diabody), and the nature of the detectable or functional label attached to the antibody. Where a radio nuclide is used for therapy, a suitable maximum single dose is about 60 mCi/m^2, to a maximum of about 250 mCi/m^2. A typical antibody dose for either tumour imaging or tumour treatment will be in the range of from 0.5 to 40 mg, preferably from 1 to 4 mg of antibody in $F(ab')_2$ form. This is a dose for a single treatment of an adult patient, which may be proportionally adjusted for children and infants, and also adjusted for other antibody formats in proportion to molecular weight. Treatments may be repeated at daily, twice-weekly, weekly or monthly intervals, at the discretion of the physician.

It is presently preferred that $F(ab')_2$ antibody fragments are used for both tumour imaging and tumour treatment.

Specific binding members of the present invention will usually be administered in the form of a pharmaceutical composition, which may comprise at least one component in addition to the specific binding member.

Thus pharmaceutical compositions according to the present invention, and for use in accordance with the present invention, may comprise, in addition to active ingredient, a pharmaceutically acceptable excipient, carrier, buffer, stabiliser or other materials well known to those skilled in the art. Such materials should be non-toxic and should not interfere with the efficacy of the active ingredient. The precise nature of the carrier or other material will depend on the route of administration, which may be oral, or by

injection, e.g. intravenous.

Pharmaceutical compositions for oral administration may be in tablet, capsule, powder or liquid form. A tablet may comprise a solid carrier such as gelatin or an adjuvant. Liquid pharmaceutical compositions generally comprise a liquid carrier such as water, petroleum, animal or vegetable oils, mineral oil or synthetic oil. Physiological saline solution, dextrose or other saccharide solution or glycols such as ethylene glycol, propylene glycol or polyethylene glycol may be included.

For intravenous, injection, or injection at the site of affliction, the active ingredient will be in the form of a parenterally acceptable aqueous solution which is pyrogen-free and has suitable pH, isotonicity and stability. Those of relevant skill in the art are well able to prepare suitable solutions using, for example, isotonic vehicles such as Sodium Chloride Injection, Ringer's Injection, Lactated Ringer's Injection. Preservatives, stabilisers, buffers, antioxidants and/or other additives may be included, as required.

A composition may be administered alone or in combination with other treatments, either simultaneously or sequentially dependent upon the condition to be treated. Other treatments may include the administration of suitable doses of pain relief drugs such as non-steroidal anti-inflammatory drugs (e.g. aspirin, paracetamol, ibuprofen or ketoprofen) or opitates such as morphine, or anti-emetics.

The present invention further provides an isolated nucleic acid encoding a specific binding member of the present invention. Nucleic acid includes DNA and RNA. In a preferred aspect, the present invention provides a nucleic acid which codes for a polypeptide of the invention as defined above, including a polypeptide as set out as residues 99 to 106 of SEQ ID NO:2 or

88 to 98 of SEQ ID NO:4, and more preferably for the entire polypeptides of SEQ ID NO:2 and SEQ ID NO:4.

The present invention also provides constructs in the form of plasmids, vectors, transcription or expression cassettes which comprise least one polynucleotide as above.

The present invention also provides a recombinant host cell which comprises one or more constructs as above. A nucleic acid encoding any specific binding member as provided itself forms an aspect of the present invention, as does a method of production of the specific binding member which method comprises expression from encoding nucleic acid therefor. Expression may conveniently be achieved by culturing under appropriate conditions recombinant host cells containing the nucleic acid. Following production by expression a specific binding member may be isolated and/or purified using any suitable technique, then used as appropriate.

Specific binding members and encoding nucleic acid molecules and vectors according to the present invention may be provided isolated and/or purified, e.g. from their natural environment, in substantially pure or homogeneous form, or, in the case of nucleic acid, free or substantially free of nucleic acid or genes origin other than the sequence encoding a polypeptide with the required function. Nucleic acid according to the present invention may comprise DNA or RNA and may be wholly or partially synthetic.

Systems for cloning and expression of a polypeptide in a variety of different host cells are well known. Suitable host cells include bacteria, mammalian cells, yeast and baculovirus systems. Mammalian cell lines available in the art for expression of a heterologous polypeptide include Chinese hamster ovary cells, HeLa cells, baby hamster kidney cells, NSO

mouse melanoma cells and many others. A common, preferred bacterial host is *E. coli*.

The expression of antibodies and antibody fragments in prokaryotic cells such as *E. coli* is well established in the art. For a review, see for example Plückthun, A. Bio/Technology **9**: 545-551 (1991). Expression in eukaryotic cells in culture is also available to those skilled in the art as an option for production of a specific binding member, see for recent reviews, for example Reff, M.E. (1993) Curr. Opinion Biotech. **4**: 573-576; Trill J.J. et al. (1995) Curr. Opinion Biotech **6**: 553-560.

Suitable vectors can be chosen or constructed, containing appropriate regulatory sequences, including promoter sequences, terminator sequences, polyadenylation sequences, enhancer sequences, marker genes and other sequences as appropriate. Vectors may be plasmids, viral e.g. 'phage, or phagemid, as appropriate. For further details see, for example, *Molecular Cloning: a Laboratory Manual*: 2nd edition, Sambrook et al., 1989, Cold Spring Harbor Laboratory Press. Many known techniques and protocols for manipulation of nucleic acid, for example in preparation of nucleic acid constructs, mutagenesis, sequencing, introduction of DNA into cells and gene expression, and analysis of proteins, are described in detail in *Short Protocols in Molecular Biology*, Second Edition, Ausubel et al. eds., John Wiley & Sons, 1992. The disclosures of Sambrook et al. and Ausubel et al. are incorporated herein by reference.

Thus, a further aspect of the present invention provides a host cell containing nucleic acid as disclosed herein. A still further aspect provides a method comprising introducing such nucleic acid into a host cell. The introduction may employ any available technique. For eukaryotic cells, suitable techniques may include calcium phosphate transfection, DEAE-Dextran,

electroporation, liposome-mediated transfection and transduction using retrovirus or other virus, e.g. vaccinia or, for insect cells, baculovirus. For bacterial cells, suitable techniques may include calcium chloride transformation, electroporation and transfection using bacteriophage.

The introduction may be followed by causing or allowing expression from the nucleic acid, e.g. by culturing host cells under conditions for expression of the gene.

In one embodiment, the nucleic acid of the invention is integrated into the genome (e.g. chromosome) of the host cell. Integration may be promoted by inclusion of sequences which promote recombination with the genome, in accordance with standard techniques.

The present invention also provides a method which comprises using a construct as stated above in an expression system in order to express a specific binding member or polypeptide as above.

The following examples illustrate the present invention.

Example 1: Isolation of Antibodies Equivalent To TNT-1

1a. Preparation of Antigen

In order to generate clones specific for human nuclear antigens, the human Burkitt's lymphoma cell line, Raji, was used as a source of antigen, prepared largely as described by G.K. Miller et. al. 1993 Hybridoma vol.12, no.6 p.p. 689-697.

Cultures were grown in RPMI-1640 + L-glutamine medium (Gibco BRL) supplemented with penicillin , streptomycin (Gibco BRL) and 5 % foetal calf serum (FetalClone II, Hyclone Europe Ltd). Cells were grown in roller bottles and produced in 1 l batches. The culture was centrifuged at 1000 rpm for 10 min and the

cells washed in PBS solution (Oxoid). The culture was spun again and the resultant pellet frozen down in 90 % FCS/ 10% DMSO. Each 1 l batch gave 1.3 - 9.5 x 10^8 cells.

To prepare nuclear extract, a cell pellet was first thawed on ice and gently resuspended in 10 ml 10 mM $CaCl_2$, 2 μM PIPES buffer. Cells were pelleted at 1000 rpm and resuspended in the same buffer containing 1 % Nonidet P40. This was incubated on ice for 10 min to allow the disruption of the cell membrane. A small aliquot was removed and analysed by light microscopy. This indicated that no intact cells remained. Nuclei were pelleted by momentary centrifugation at 3000 rpm. The nuclei (pearly white pellet) were resuspended in 5 ml CaPIPES buffer. 1 ml nuclei were pelleted and resuspended in 1 ml sterile TE pH8.0. This was subjected to 3 x 45 sec sonication bursts to disrupt the nuclear membrane. Debris was pelleted at 13 000 rpm (microfuge) for 2 min and the supernatant transferred to a fresh tube. The amount of nuclear extract was determined by absorption at 260 nm using the conversion factor 10A260 = 1 mg nucleohistone. The yield of nucleohistone was estimated to be 1.2 mg/ml.

1b. Induction of phage antibody libraries

Two different phage antibody repertoires were selected for antibodies to Raji cell nuclear extract, the VH synthetic (Nissim et al., 1994) and large scFv' (Vaughan et al 1996) repertoires were each treated as follows in order to rescue phagemid particles. 500 ml prewarmed (37°C) 2YTAG (2YT media supplemented with 100 μg/ml ampicillin and 2 % glucose) in a 2 l conical flask was inoculated with approximately 3 x 10^{10} cells from a glycerol stock (-70°C) culture of the appropriate library. The culture was grown at 37°C with good aeration until the OD600nm reached 0.7 (approximately 2 hours). M13K07 helper phage (Stratagene) was added to the culture to a multiplicity of infection (moi) of approximately 10 (assuming

that an OD600nm of 1 is equivalent to 5 x 10^8 cells per ml of culture). The culture was incubated stationary at 37°C for 15 minutes followed by 45 minutes with light aeration (200 rpm) at the same temperature. The culture was centrifuged and the supernatant drained from the cell pellet. The cells were resuspended in 500 ml 2YTAK (2YT media supplemented with 100 μg/ml ampicillin and 50 μg/ml kanamycin), and the culture incubated overnight at 30°C with good aeration (300 rpm). Phage particles were purified and concentrated by a polyethylene glycol (PEG) precipitations (Sambrook et al., 1990) and resuspended in 10 ml 1 x TE. 0.5 g/ml caesium chloride was added to each dissolved, transferred to ultracentrifuge tubes and centrifuged at 42 000 rpm at 16°C in a Sorvall TFT 65.13 rotor overnight. The caesium banded phage were removed with a needle and syringe and dialysed overnight in 2 x 2l TE at 4°C using a Slide-a Lyzer Cassette (Pierce) according to the manufacturers instructions. Phage were stored at 4°C until use.

1c. Panning of phage antibody library on Nuclear Extract

Phage induced from the two repertoires were each separately panned on Raji cell nuclear extract, the same antigen as that used for the inoculation of mice to give TNT-1. For the large scFv repertoire, 1 x 10^9 sub-aliquots of the library were selected separately. A 75mm x 12mm immuno tube (Nunc; Maxisorp) was coated with 10 μg nuclear extract in 1 ml sterile PBS overnight at 4°C, followed by 1 hour at 37°C. After washing 3 times with PBS, the tube was filled with 3%MPBS (3 % 'Marvel' skimmed milk powder, 1x PBS) and incubated for 1 hour at 37°C for blocking. The wash was repeated, phagemid particles (1012 tu) in 2 ml of 3% MPBS were added and the tube incubated stationary at 37°C for 1 hour. The tube was washed 10 times with PBST(0.1%), then 10 times with PBS. Bound phage particles were eluted from the tube by adding 1 ml of 100mM-triethylamine, and incubating the tube stationary at room

temperature for 10 minutes. The eluted material was immediately neutralised by pipetting into a tube containing 1 ml 1M-Tris.HCl (pH7.4). Phage were stored at 4°C. 1.5 ml of eluted phage were used to infect 10 ml of logarithmically growing *E. coli* TG1 (Gibson, 1984). Infected cells were grown for 1 hour at 37°C with light aeration in 2YT broth, and then plated on 2YTAG medium in 243mm x 243mm dishes (Nunc). Plates were incubated overnight at 30°C. Colonies were scraped off the plates into 10 ml of 2YT broth and 15 % (v/v) glycerol added for storage at -70°C.

Glycerol stock cultures from the first round of panning of each of the two repertoires on nuclear extract were rescued using helper phage to derive phagemid particles for the second round of panning. 250 µl of glycerol stock was used to inoculate 50 ml 2YTAG broth, and incubated in a 250 ml conical flask at 37°C with good aeration until the OD600nm reached 0.7 (approximately 2 hours). M13K07 helper phage (moi=10) was added to the culture which was then incubated stationary at 37°C for 15 minutes followed by 45 minutes with light aeration (200 rpm) at the same temperature. The culture was centrifuged and the supernatant drained from the cell pellet. The cells were resuspended in 50 ml prewarmed 2YTAK, and the culture incubated overnight at 30°C with good aeration.

Phage induced from the first round of panning of each of the two repertoires, was selected a second time as described above . After extensive washing, bound phage were eluted from the tube using 1 ml of 100 mM-triethylamine, neutralised by the addition of 0.5 ml 1M-Tris.HCl (pH7.4) and infected into TG1 cells as before.

1d. Growth of single selected clones for immunoassay

Individual colonies from the second round selection were used to inoculate 100 µl 2YTAG into individual wells of 96 well

tissue culture plates (Corning). Plates were incubated at 30°C overnight with moderate shaking (200 rpm). Glycerol to 15 % was added to each well and these master plates stored at -70°C until ready for analysis.

1e. ELISA to Identify Nuclear Extract Binding scFv

Clones specific for nuclear extract were identified by ELISA, using scFv displayed on phage. Cells from the master plates were used to inoculate fresh 96 well tissue culture plates containing 100 µl 2YTAG per well. These plates were incubated at 37°C for 6-8 hours or until the cells in the wells were growing logarithmically (OD600 0.2-1.0). M13K07 was added to each well to an moi of 10 and incubated stationary for 15 min then 45 min with gentle shaking (100 rpm), both at 37°C. The plates were centrifuged at 2000 rpm for 10 min and the supernatant eluted. Each cell pellet was resuspended in 100 µl 2YTAK and incubated at 30°C overnight.

Each plate was centrifuged at 2000 rpm and the 100 µl supernatant from each well recovered and blocked in 20 µl 18%M6PBS (18 % skimmed milk powder, 6 x PBS), stationary at room temperature for 1 hour. Meanwhile, flexible microtitre plates which had been blocked overnight stationary at 4°C with either 50 µl 2.5 µg/ml nuclear extract in PBS or 50 µl PBS alone (giving an uncoated control plate), were washed 3 times in PBS and blocked for 2 h stationary at 37°C in 3MPBS. These plates were then washed three times with PBS and 50 µl preblocked phage added to each well of both the nuclear extract-coated or uncoated plate. The plates were incubated stationary at 37°C for 1 h after which the phage were poured off. The plates were washed by incubating for 2 min in PBST three times followed by incubating for 2 min in PBS three times, all at room temperature.

To each well of both the nuclear extract-coated and the

uncoated plate, 50 µl of a 1 in 10 000 dilution of sheep anti-fd antibody (Pharmacia) in 3MPBS was added and the plates incubated at 37°C stationary for 1 h. Each plate was washed as described above and 50 µl of a 1 in 5 000 dilution donkey anti-sheep alkaline phosphatase conjugate (Sigma) in 3MPBS added and incubated stationary at 37°C for 1 h. Plates were washed as described as above followed by two rinses in 0.9% NaCl. Alkaline phosphatase activity was visualised using the chromagenic substrate pNPP (Sigma). The absorbance signal generated by each clone was assessed by measuring the optical density at 405 nm using a microtitre plate reader.

Clones were chosen for further analysis if the ELISA signal generated on the nuclear extract-coated plate was at least double that on the uncoated plate. About 350 out of 700 clones analysed met the criteria set. Eighty six of these clones were selected for further characterisation.

1f. Specificity ELISA

Eighty six clones, identified as binding nuclear extract rather an uncoated well, were further analysed for fine specificity. Specificity ELISA's were carried out using scFv displayed on phage as described above. Microtitre plate wells were coated with 50 µl of either 5 µg/ml nuclear extract, 5 µg/ml histone H1 (Boehringer Mannheim), 100 µg/ml lysozyme or PBS (the uncoated well) at 4°C overnight in PBS then 1 hour at 37°C. After preblocking both the phage and the microtitre plates, 50 µl blocked phage from each clone was added to a well coated with each antigen or an uncoated well. As above, alkaline phosphatase activity was visualised using either the chromagenic substrate pNPP (Sigma). Clones which were specific for nuclear extract and histone H1 but showed no cross-reaction with lysozyme or the uncoated well were analysed further. Clones were designated specific for nuclear extract and histone H1 if the ELISA signal generated on both antigens was at least

five-fold greater than the signal on either lysozyme or on an uncoated well. TNT-1 binds both nuclear extract and histone H1 but not lysozyme. Twenty nine of the clones gave a binding profile similar to TNT-1 and were characterised further.

Example 2: Western Blot Analysis

2a. Preparation of soluble scFv

The twenty nine clones with a binding profile similar to TNT-1 were analysed for binding to nuclear extract components by Western blot analysis. Soluble scFv prepared from the periplasm of 50 ml culture was used for this. 2 ml overnight culture (grown at 30°C in 2TYAG) was added to 50 ml 2TYA and incubated at 30°C to OD600 0.6. IPTG was added to a final concentration of 1 mM and the culture grown overnight at 30°C. Cells were pelleted in 50 ml Falcon tubes and taken up in 5 ml ice-cold 50 mM Tris-Cl pH8, 20% Sucrose, 1mM EDTA. The suspension was incubated on ice for 30 min then pellet at 8K for 20 min in a SM-24 rotor. The supernatant was used for the Western blot analysis.

2b. Casting of SDS-Polyacrylamide Gel.

An SDS-Polyacrylamide gel was mounted in a Bio-Rad electrophoresis cell and 100 ml 2.5 x SDS loading buffer added into the slot to create a spirit level surface. 200 ml nuclear extract sample was heated for 2 min at 90°C in 1 x SDS loading buffer with betamercaptoethanol and applied to the top of the 2.5 x SDS loading buffer. The gel was run at 20 mA until the bromophenol blue dye front had passed through the bottom of the gel. The protein was blotted from the gel on to a PVDF Immobilon-P transfer membrane, using a Bio-Rad semi-dry transfer cell 24 V. The membrane was briefly washed in methanol and air dried. The membrane was sliced into approximately 3 mm strips.

2c. Western blotting

The membrane strips from above were briefly washed in methanol then water. Each membrane strip was blocked in 5 % Marvel/PBS/Tween for 45 min in a Costar reagent reservoir divided in to 12 individual wells. Periplasmic extract of each clone was preblocked in Marvel/PBS by adding 0.5 ml 18 % Marvel, 6 x PBS to 2.5 ml of periplasmic preparation and incubating for 15 min at room temperature. The preblocked antibodies were added to each strip and incubated at room temperature for 1 h. The strips were washed for 30 min with several changes of PBS/Tween then transferred to another reagent reservoir and washed with PBS 2 x 5 min. 1/100 diluted 9E10 anti-myc tag in 3 % Marvel/PBS was added to the strips and incubated at room temperature for 1 h. The strips were washed for 30 min with several of changes PBS/Tween then transferred to another reagent reservoir and washed with PBS 2 x 5 min. 1/1000 diluted goat anti-mouse IgG peroxidase in 3 % Marvel/PBS was added to the strips and incubated at room temperature for 1 h. The strips was washed for 1 h with several of changes PBS/Tween then transferred to another reagent reservoir and washed with PBS 2 x 5 min and saline 2 x 5 min. Each strip was washed briefly in water, transferred to a new reagent reservoir and developed using ECL (Amersham) according to the manufacturers instruction. The film was exposed to the strips for between 2 min and 16 h.

A second western blot of selected candidates on bovine histone H1 preps and nuclear extract was performed using a 20 % PhastGel using the Pharmacia Phast Gel system essentially as above.

The western blot showed that a large number of the candidates had a binding profile close to the binding profile of TNT1, recognising histone H1 and histone H3 but also binding to some extent to the other core histones and to other non-identified proteins in the preparation. Seven clones had almost identical

binding by western blot analysis as TNT-1, NHS 16, NHS 19, NHS 39, NHS 45, NHS 65, NHS 76 and NHS 86. These TNT-1 equivalent clones were chosen for further analysis. A further clone, NHS 22 was also chosen for further analysis, it has a similar ELISA profile to the other candidates but a different Western blot pattern.

Not all candidates showed a TNT1 like profile, some clones did not give a sufficient signal to determine any binding profile and other clones had a Western blot profile that was different from the TNT1 profile. It is infact surprising that so many of the clones have similar profiles to the TNT1 Western blot profile taking in to account the complexity of the starting antigen.

2d. Sequencing of TNT-1 Equivalent ScFv Antibodies

The nucleotide sequence of the TNT-1 equivalent human scFv antibodies were determined by first using vector-specific primers to amplify the inserted DNA from each clone. Cells from an individual colony on a 2YTAG agar plate were used as the template for a polymerase chain reaction (PCR) amplification of the inserted DNA using the primers pUC19 reverse and fdtet sequence. pUC19 reverse has the sequence: 5' AGC GGA TAA CAA TTT CAC ACA GG 3' (SEQ ID NO:5). fdtet sequence has the sequence: 5' GTC GTC TTT CCA GAC GTT AGT 3' (SEQ ID NO:6). Amplification conditions consisted of 30 cycles of 94°C for 1 min, 55°C for 1 min and 72°C for 2 min, followed by 10 min at 72°C. The PCR products were purified using a PCR Clean-up Kit (Promega) in to a final volume of 50 μl H_20. Between 2 and 5 μl of each insert preparation was used as the template for sequencing using the Taq Dye-terminator cycle sequencing system (Applied Biosystems). The primers mycseq10 and PCR-L-Link were used to sequence the light chain of each clone and PCR-H-Link and pUC19reverse to sequence the heavy chain. The sequences of these primers are as follows:

pUC19 reverse 5' AGC GGA TAA CAA TTT CAC ACA GG 3' (SEQ ID NO: 7)

myc sequence 10 5' CTC TTC TGA GAT GAG TTT TTG 3' (SEQ ID NO: 8)

PCR-H-Link 5' ACC GCC AGA GCC ACC TCC GCC 3' (SEQ ID NO: 9)

PCR-L-Link 5' GGC GGA GGT GGC TCT GGC GGT 3' (SEQ ID NO: 10)

2e. Sequence and Source of the TNT-1 Equivalent ScFv Antibodies

Eight different TNT-1 equivalent antibodies were isolated from the selections using the two libraries described above. A further clone, D3, was isolated by the epitope imprinting of chTNT-1 using nuclear extract as the source of antigen, much as described in Jespers *et al*, (1994), Biotechnology 12; 899-903. Each clone name, its origin and its heavy and light chain germline is given below. The sequences of NHS 76 VH and VL chains (SEQ ID NOs: 2 and 4) are shown in Figures 1 and 2.

CLONE	LIBRARY SOURCE	VH GERMLINE	VL GERMLINE
NHS 16	large scFv	DP 71	DPL 16
NHS 19	large scFv	DP 75	DPL 11
NHS 22	large scFv	DP 47	L12a
NHS 39	large scFv	DP 75/15	DPL 11
NHS 45	large scFv	DP 46	L12a
NHS 65	large scFv	DP 7	DPL 11
NHS 76	synthetic	DP 67	DPL 16
NHS 86	large scFv	DP 47	L12 a

There does not appear to be any correlation between the specificity of these eight clones and their sequence. A wide range of VH/VL combinations are utilised and the CDR regions are highly variable. It is therefore unlikely that antibodies with equivalent in vitro binding characteristics to TNT-1 could be predicted from their amino acid sequence.

Example 3: Conversion of Candidates to Whole Antibody Format.

Restriction digests, ligations, clonings and PCRs

Restriction digests, ligations and cloning of DNAs were essentially as described (Sambrook et al., 1987).
Each PCR of 50 µl contained 10 mM Tris HCl pH 8.85, 25 mM KCl, 5 mM $(NH_4)_2SO_4$, 2 mM $MgSO_4$, 250 µM dNTPs, 0.5 µM each primer, 1U PwoI DNA polymerase (Boehringer Mannheim) and appropriate amounts of template DNA. Each reaction mixture was subjected to 15-25 cycles of 94°C, 30 s; 50°C, 30 s; 72°C, 60s.

Construction of cell line expressing NHS76 IgG

For the construction of cell lines expressing human IgG1 antibodies the heavy and light chain variable domains from the NHS76 scFv expressing phage were cloned into mammalian

expression vectors containing human IgG1 and human lambda constant domains respectively.

To facilitate possible subsequent cloning of the VH and VL domains of NHS76 into a single expression vector necessitated deletion of internal BamHI sites. The restriction sites were removed by PCR-directed mutagenesis (Ho, S.N., Hunt, H.D., Horton, R.M., Pullen, J.K. and Pease, L.R. 1989 Gene 77, 51-59) using primers P54 (SEQ ID NO: 15) + P58 (SEQ ID NO: 17) and P57 (SEQ ID NO: 16)+ P17 (SEQ ID NO: 13) for NHS76 VH, and primers P61 (SEQ ID NO: 20) + P60 (SEQ ID NO: 19) and P59 (SEQ ID NO: 18) + P45 (SEQ ID NO: 14) for NHS76 Vλ. All oligonucleotide primers of Example 3 are shown in Table 6. The PCR products of the expected sizes were joined in a second PCR using P54 + P17 for VH, and P61 + P45 for Vl. No changes in amino acid sequences resulted with the removal of the restriction sites.

Heavy chain expression vector The NHS76 VH DNA amplified with P54 and P17 was joined by overlapping PCR to a 158 bp DNA fragment containing a signal sequence, splice sites and intron from M13VHPCR1 (Orlandi et al., 1989) using oligonucleotides P10 (SEQ ID NO: 11) and P17. The 539 bp PCR product was cut with HindIII and ApaI and cloned into HindIII-ApaI cut pGamma1 (obtained from Lonza). Ligated DNA was transformed into E. coli TG1 and ampicillin-resistant colonies screened. A plasmid with the correct insertion was identified and designated pNHS76γ1.

Light chain expression vector The vector for the expression of lambda chains was a modified version of pMR15.1 (obtained from Lonza) where the Cκ DNA was replaced with human Cλ DNA and was designated pCλ. The NHS76 Vλ DNA amplified with P61 and P45 was joined by overlapping PCR to a 168 bp DNA fragment containing a signal sequence, splice sites and intron from M13VKPCR1 (Orlandi et al., 1989) using oligonucleotides P11 (SEQ ID NO: 12) and P45. The 529 bp PCR product was cut with BstBI and PacI and cloned into BstBI-PacI cut pCλ. Ligated DNA was transformed into E. coli TG1 and ampicillin-resistant colonies screened. A plasmid with the correct insertion was

identified and designated pNHS76l.

NHS76 IgG Expression

NHS76 IgG was expressed in the mouse myeloma cell line NS0 (ECACC 85110503). 45 μg of pNHS76γ1 and 15 μg pNHS76λ DNAs were linearised by digestion with PvuI, ethanol precipitated and dissolved in 100 μl water. 10^7 NS0 cells were washed in PBS, resuspended in 0.9 ml PBS, mixed with the vector DNA and held in ice for 5 min. The cells were then electroporated with 2 pulses of 1.5 kV at 3 μFd and incubated in ice for 10 min. The transfected cells were then added to 30 ml Dulbecco's modified Eagle's medium (DMEM) containing 2 mM glutamine and 10 % dialysed foetal calf serum (FCS) as described by Bebbington et al. (1992) and 50 μl aliquots distributed into 6 x 96-well plates. 24 h later glutamine-free DMEM / 10 % FCS (Bebbington et al. 1992) was added to each well. Three to 6 weeks after transfection colonies were screened by ELISA for the ability to secrete human IgG. Wells of ELISA plates (Immulon 4, Dynatech) were coated in 50 mM sodium bicarbonate / carbonate pH 9.6 with 100 ng per well of goat anti-human IgG antibodies (Sera-Lab). Supernatant from wells containing transfected colonies was added to the wells in PBS containing 0.05 % (v/v) Tween 20 (PBST) for 1 h. The plates were washed 3 times with PBST and captured human IgG was detected with 100 μl 1:2000 - 1:5000 dilution horseradish peroxidase (HRP) conjugated goat anti-human lambda antibodies in PBST (Sera-Lab). After 30 min at room temperature the plates were washed 3 x PBST and 100 μl OPD substrate added (50μl). Reactions were stopped after 5-10 min by the addition of 50 μl 12.5 % (v/v) sulphuric acid and the A 490 nm measured. Transfectants secreting the highest amounts of IgG were expanded for growth in glutamine-free medium in reduced FCS, in gammaglobulin-free FCS or no FCS. Cell lines were subsequently cloned by limiting dilution.

Purification of antibodies

Human IgG1 antibodies were purified by protein A affinity chromatography. Supernatant from the growth of transfected NS0 cells secreting NHS76 IgG was clarified by centrifugation and passage through a 0.22 μm filter. A protein A Sepharose column (HiTrap, Pharmacia) was equilibrated with PBS and the supernatant applied. The column was then washed with 10 column volumes of PBS and any bovine IgG present (from the foetal calf serum in the culture medium) removed by elution with 0.1 M sodium acetate pH 4.5. After further washing with PBS human IgG was eluted with 0.1 M sodium acetate pH 3.5. Eluted fractions were neutralised with 1 M Tris HCl pH 9.0 and protein containing fractions identified by measuring the absorbance at 280 nm. Antibody containing fractions were pooled and dialysed against PBS.

Example 4: Digestion of Whole Antibodies

For the animal model, the candidates were tested as both whole antibodies and as F(ab')2 fragments. These were generated by limited pepsin digestion. A trial digest was set up for each antibody tested. The following were added to a 1.5 ml microfuge tube:

-25 μl whole antibody in PBS

-5 μl 1M sodium acetate buffer at either pH 3, 3.5, 4, 4.5, 5 or 5.5

-pepsin at a final concentration of 10 μg pepsin / mg antibody (enzyme/substrate = 1/100).

- water to give a total volume of 50 μl

The digest was incubated at 37°C and samples taken out after 1, 4, 8 and 16 hours. The reaction was stopped by adding 3M Tris-HCl pH8.8 to a final concentration of 0.3 M and analysed on a 10-15 % Phast gel (Pharmacia).

This gave the optimum digestion conditions for each antibody and was applied to the large-scale digests. Following digestion, the F(ab')2 fragment was purified on a 24 ml S200

FPLC column and eluted in the 14-16.5 ml fractions. These were pooled and concentrated to about 4 mg/ml in an Amicon Centriprep YM10.

Example 5: Iodination of Antibodies

For the iodination, first the required number of screw top eppendorf tubes were coated with iodogen. Iodogen (Pierce) was dissolved at 1.2mg/ml in chloroform, then a 1 in 10 dilution in chloroform made. 0.5 ml was added to each tube and carefully blown off the under N2. These could be stored in a desiccator at 4°C for future use. To iodinate, 0.5 mg of protein was transferred to an iodogen tube on ice and made up to 0.5 ml with borate labelling buffer (100 ml 100 mM Borax, 100 ml 0.9% NaCl, pH to 8.5 with HCl; Filter Sterilised). An appropriate volume of Na125I (250 μCi - 2.5 mCi) was added, mixed and left on ice for 10 minutes. Using a fine pasteur pipette, the whole reaction was transferred to the top of a PD-10 column (Pharmacia Biotech) which has been pre-blocked with 5% BSA/PBS and washed through 5 times with sterile PBS. 0.5 ml aliquots of sterile PBS were added sequentially to the top of the column and run through. Similar 0.5 ml aliquots were collected at the bottom of the tube. Each fraction was then be counted in a gamma counter. The iodinated protein usually comes out in fractions 6-7.

Example 6: Biodistribution studies.

Experiments were conducted to establish the ability of $F(ab')_2$ fragments of NHS76 to localise to human tumour xenografts in athymic mice. Female athymic nu/nu MF-1 mice (4 per group) were injected subcutaneously with 10^7 ME-180 human cervical carcinoma cells (ATCC HTB-33). Approximately 3-4 weeks later tumour-bearing mice were injected in the tail vein with 10 μg (37 kBq) ^{125}I-labelled NHS76 IgG or ^{125}I-labelled NHS76 $F(ab')_2$. At selected time points up to 96 h post-injection mice were sacrificed and blood, tumour and all major organs were

collected and weighed (4 mice per time point). Radioactivity was measured in a gamma counter (Cobra II, Packard). The percentage of the injected dose per gram of each organ was determined and the organ : blood ratios calculated. The ME-180 xenograft model is used in the art as an indicator for human clinical utility of test substances. See Epstein *et al* in "Handbook of Targeted Delivery of Imaging Agents", *ibid*. Table 1 and Fig. 3 show the biodistribution of ^{125}I-labelled NHS76 F(ab')2. Fig. 3a shows the distribution, and Fig. 3b the mean tissue: blood ratio. The fragment is rapidly cleared from the circulation with only 1 % of the injected dose remaining at 24 h p.i. NHS76 F(ab')2 demonstrates increased tumour localisation with time such that at 96 h p.i the tumour : blood ratio is 4.3 : 1. There is no specific retention in any other organ examined.

Figs. 4a and 4b show the biodistribution and mean tumour:blood ratio respectively of 125I-labelled D3 F(ab')2. The data are shown in Table 2. D3 demonstrates a more rapid clearance from the circulation than NHS76 F(ab')2 with only 0.29 % of the injected dose remaining in the circulation at 24 h. D3 F(ab')2 also shows specific tumour localisation and at 96 h shows a tumour : blood ratio of 5.6 : 1. There is also, however, retention of 125I in the skin (skin : blood ratio of 3.3 :1). In comparison to NHS76, D3 demonstrates a much lower amount of accretion in tumour, for example at 96 h p.i. about there is about 6 x less radioactivity in the tumour with D3 F(ab')2 than NHS76 F(ab')2.

Tables 3 and 4 show the biodistribution of 125I-labelled F(ab')2 fragments of NHS45 and NHS86. Both fragments are cleared from the circulation more rapidly than NHS76 F(ab')2 with 0.1 % of the injected dose remaining at 24 h p.i. From 24 h to 96 h p.i. both antibody fragments show uptake to tumour and exhibit tumour : blood ratios of > 3-5. However, fragments

of both NHS45 and NHS86 also show significant retention in other organs.

NHS19 F(ab')2 shows little, if any, specific tumour uptake (Table 5, see also Figs. 5a and 5b).

Thus NHS76 F(ab')2,in contrast to the fragments of the other antibodies, shows the ability to localise to tumour with no specific retention in any other organ. The antigen binding specificity, the serum clearance and its entirely human makeup gives NHS76 F(ab')2 a composite of properties well suited for use in humans as an in vivo diagnostic or therapeutic agent with the potential for repeated administration.

TABLES 1-5 Biodistribution of 125I-labelled F(ab')2 fragments in ME-180 tumour-bearing athymic mice. Values represent the mean % injected dose (ID)/ g; (SD of organs from 4-5 mice).

TABLE 1 - NHS76 F(ab')2 : % ID / g TISSUE										
ORGAN	Blood	Colon	Femur	Kidney	Liver	Lung	Muscle	Skin	Spleen	Tumour
6 h	7.96 (0.38)	1.95 (0.18)	1.41 (0.20)	9.08 (0.92)	3.97 (0.27)	3.89 (0.24)	0.73 (0.13)	2.59 (0.40)	3.47 (0.18)	1.99 (0.40)
24 h	1.06 (0.27)	0.39 (0.11)	0.39 (0.07)	1.31 (0.33)	0.51 (0.11)	0.79 (0.18)	0.20 (0.05)	0.64 (0.20)	0.52 (0.08)	1.07 (0.24)
48 h	0.20 (0.04)	0.07 (0.01)	0.08 (0.02)	0.32 (0.05)	0.11 (0.02)	0.20 (0.03)	0.04 (0.01)	0.15 (0.02)	0.10 (0.02)	0.38 (0.05)
96 h	0.07 (0.01)	0.02 (0.003)	0.08 (0.02)	0.14 (0.02)	0.04 (0.01)	0.07 (0.02)	0.014 (0.001)	0.06 (0.005)	0.05 (0.01)	0.29 (0.24

TABLE 2 - D3 F(ab')2 : % ID / g TISSUE										
ORGAN	Blood	Colon	Femur	Kidney	Liver	Lung	Muscle	Skin	Spleen	Tumour
6 h	3.51 (0.23)	0.92 (0.07)	0.61 (0.09)	12.6 (3.2)	1.06 (0.06)	2.24 (0.82)	0.48 (0.17)	1.13 (0.25)	0.94 (0.10)	1.24 (0.40)
24 h	0.29 (0.05)	0.09 (0.01)	0.08 (0.01)	0.72 (0.12)	0.10 (0.01)	0.22 (0.02)	0.07 (0.02)	0.22 (0.05)	0.10 (0.01)	0.49 (0.21)
48 h	0.04 (0.02)	0.02 (0.01)	0.02 (0.005)	0.10 (0.05)	0.02 (0.01)	0.04 (0.02)	0.01 (0.003)	0.06 (0.01)	0.02 (0.01)	0.14 (0.11)
96 h	0.008 (0.002)	0.007 (0.002)	0.007 (0.002)	0.03 (0.01)	0.006 (0.002)	0.01 (0.004)	0.005 (0.001)	0.03 (0.006)	0.01 (0.003)	0.05 (0.04)

TABLE 5 - NHS19 F(ab')2 : % ID / g TISSUE										
ORGAN	Blood	Colon	Femur	Kidney	Liver	Lung	Muscle	Skin	Spleen	Tumour
6 h	0.74 (0.16)	0.41 (0.22)	0.33 (0.26)	1.36 (0.21)	0.40 (0.08)	0.53 (0.15)	0.19 (0.13)	0.63 (0.29)	0.35 (0.10)	0.51 (0.11)
24 h	0.11 (0.01)	0.05 (0.01)	0.07 (0.07)	0.43 (0.13)	0.07 (0.003)	0.10 (0.015)	0.03 (0.02)	0.12 (0.02)	0.06 (0.01)	0.16 (0.04)
48 h	0.07 (0.03)	0.04 (0.01)	0.03 (0.008)	0.25 (0.07)	0.05 (0.02)	0.06 (0.02)	0.02 (0.003)	0.09 (0.04)	0.04 (0.01)	0.11 (0.04)
96 h	0.035 (0.019)	0.014 (0.005)	0.017 (0.005)	0.097 (0.061)	0.023 (0.014)	0.030 (0.014)	0.009 (0.003)	0.048 (0.015)	0.021 (0.010)	0.053 (0.019)

TABLE 3 - NHS45 F(ab')2 : % ID / g TISSUE

ORGAN	Blood	Colon	Femur	Kidney	Liver	Lung	Muscle	Oesophagus	Skin	Small Intestine	Spleen	Stomach	Tumour
6 h	0.71 (0.42)	0.34 (0.22)	0.42 (0.25)	2.82 (1.57)	0.54 (0.34)	0.81 (0.58)	0.30 (0.19)	0.68 (0.43)	0.61 (0.38)	0.47 (0.26)	0.50 (0.28)	1.20 (0.66)	0.72 (0.47)
24 h	0.10 (0.005)	0.08 (0.02)	0.09 (0.02)	0.30 (0.05)	0.15 (0.04)	0.15 (0.02)	0.04 (0.01)	0.19 (0.09)	0.23 (0.06)	0.10 (0.02)	0.12 (0.03)	0.19 (0.06)	0.23 (0.03)
48 h	0.027 (0.006)	0.023 (0.006)	0.037 (0.008)	0.12 (0.05)	0.064 (0.017)	0.059 (0.037)	0.011 (0.005)	0.053 (0.004)	0.050 (0.025)	0.033 (0.013)	0.074 (0.024)	0.041 (0.012)	0.15 (0.033)
96 h	0.007 (0.003)	0.012 (0.003)	0.020 (0.005)	0.028 (0.017)	0.013 (0.008)	0.016 (0.007)	0.005 (0.001)	0.049 (0.011)	0.020 (0.005)	0.019 (0.008)	0.027 (0.016)	0.011 (0.001)	0.026 (0.014)

TABLE 4 - NHS86 F(ab')2 : % ID / g TISSUE

ORGAN	Blood	Colon	Femur	Kidney	Liver	Lung	Muscle	Oesophagus	Skin	Small Intestine	Spleen	Stomach	Tumour
6 h	1.62 (0.42)	1.07 (0.22)	1.51 (0.25)	6.45 (1.57)	5.23 (0.34)	2.58 (0.58)	0.85 (0.19)	1.56 (0.43)	1.66 (0.38)	2.57 (0.26)	3.57 (0.28)	3.47 (0.66)	1.75 (0.47)
24 h	0.12 (0.01)	0.14 (0.02)	0.23 (0.02)	0.50 (0.06)	0.34 (0.04)	0.35 (0.11)	0.06 (0.01)	0.22 (0.05)	0.22 (0.02)	0.19 (0.02)	0.38 (0.02)	0.23 (0.03)	0.29 (0.05)
48 h	0.072 (0.032)	0.067 (0.027)	0.12 (0.04)	0.28 (0.10)	0.13 (0.04)	0.19 (0.09)	0.029 (0.007)	0.19 (0.11)	0.12 (0.02)	0.099 (0.029)	0.22 (0.13)	0.091 (0.041)	0.25 (0.07)
96 h	0.030 (0.009)	0.027 (0.013)	0.053 (0.030)	0.079 (0.027)	0.044 (0.019)	0.049 (0.010)	0.011 (0.002)	0.064 (0.015)	0.043 (0.012)	0.029 (0.011)	0.061 (0.016)	0.035 (0.014)	0.054 (0.012)

TABLE 6 - OLIGONUCLEOTIDE PRIMERS	
P10	5'-CTAAGCTTACTGAGCACACAGGACCTCACC-3'
P11	5'-AATTTTCGAACTACAGTTACTGAGCACACAGGACC-3'
P17	5'-ATGGGCCCTTGGTGGAAGCTGAAGAGACGGTGACCAGGGTGCC-3'
P45	5'-GCAAAGTTAATTAATTCTACTCCACCTAGGACGGTCAGCTTGGTCCCTCCGCCGAA-3'
P54	5'-TTTGGATATCTCTCCACAGGTGTCCACTCCCAGGTGCAGCTGCAGGAGTCCGGCCCA-3'
P57	5'-CTGGGGCTGGATTCGGCAGCCCCCA-3'
P58	5'-TGGGGGCTGCCGAATCCAGCCCCAG-3'
P59	5'-GCCCTCAGGGATTCCAGACCGATTC-3'
P60	5'-GAATCGGTCTGGAATCCCTGAGGGC-3'
P61	5'-TTGGATATCTCTCCACAGGTGTCCACTCCTCTTCTGAGCTGACTCAGGACCCT-3'

Example 7: Antigen specificity of NHS76

The ability of NHS76 IgG to bind to histones was examined by ELISA. Wells of microtitre plates (Immulon IV, Dynatech) were coated with 100ng bovine histones (Boeringher Mannheim) in 50 mM bicarbonate/carbonate buffer pH9.6 at 4C for 16h. Wells were blocked with 200μl 2% Marvel in PBST (MPBST) for 1h at 37C. After washing in PBST, NHS76 IgG in 100μl MPBST was added to the wells and incubated for 1h at room temperature. After washing, bound IgG was detected by the addition of 100μl 1:2000 HRP goat anti-human IgG antibodies (Harlan Sera-Lab). After 30 min at room temperature the wells were washed and 100μl o-phenylenediamine (0.4 mg/ml in 24 mM citric acid, 52 mM Na_2HPO_4 pH 5.2, 0.003% H_2O_2) added. After 6 min the reaction was stopped by the addition of 50μl 12.5% sulphuric acid and the absorbance at 490nm measured. The relative binding of the NHS76 IgG is histone H1 > histone H3 > histone H2B > histone H4 = histone H2A (Figure 6). A competition ELISA to determine the relative binding of NHS76 IgG to histones in solution was also employed and confirmed the ELISA data.

The ability of NHS76 IgG to bind histones from human and other species was analysed by Western blotting. Nuclear extracts of Raji (human), NS0 (mouse), JTC-19 (rat) and JH4 (guinea pig) cells were electrophoresed through a reducing 14% SDS-PAG. The gel was electroblotted onto nitrocellulose membrane (Amersham International) which was subsequently blocked with MPBST for 1h at room temperature. The membrane was then incubated with NHS76 IgG at 1μg/ml for 50 min at room temperature. After 4 washes with PBST the membrane was incubated with 1:5000 HRP goat anti-human IgG in MPBST for 40 min at room temperature. After extensive washing the membrane was incubated with ECL reagent (Amersham) according to the manufacturer's instructions. The western blot result (Figures 7a-c (3 different exposures to show bands at different intensities)) demonstrates the binding of NHS76 IgG to proteins from human, mouse, rat and guinea pig cells which comigrate with bovine histones H1 and H3.

CLAIMS

1. An isolated specific binding member capable of binding an intracellular antigen, wherein said specific binding member comprises a polypeptide binding domain comprising an amino acid sequence substantially as set out as residues 99 to 106 of SEQ ID NO:2.

2. An isolated specific binding member according to claim 1 which further comprises the polypeptide binding domains substantially as set out as residues 31-36 and 51-66 of SEQ ID NO:2.

3. An isolated specific binding member according to claim 2 wherein said binding domains are carried by a human antibody framework.

4. An isolated specific binding member according to claim 3 which comprises substantially the polypeptide sequence of SEQ ID NO:2.

5. An isolated specific binding member capable of binding an intracellular antigen, wherein said specific binding member comprises a polypeptide binding domain comprising an amino acid sequence substantially as set out as residues 88 to 98 of SEQ ID NO:4.

6. An isolated specific binding member according to claim 5 which further comprises the polypeptide binding domains substantially as set out as residues 23-33 and 49-55 of SEQ ID NO:4.

7. An isolated specific binding member according to claim 6 wherein said binding domains are carried by a human

antibody framework.

8. An isolated specific binding member according to claim 7 which comprises substantially the polypeptide sequence of SEQ ID NO:4.

9. A specific binding member which comprises a first specific binding member as defined in any one of claims 1 to 4 in association with a second specific binding member as defined in any one of claims 5 to 8.

10. A specific binding member according to claim 9 in the form of an antibody F(ab')2 or scFv fragment.

11. A specific binding member according to any one of claims 1 to 10 which carries a detectable or functional label.

12. An isolated nucleic acid which comprises a sequence encoding a specific binding member as defined in any one of claims 1 to 11.

13. A method of preparing a specific binding member as defined in any one of claims 1 to 11 which comprises expressing the nucleic acid of claim 12 under conditions to bring about expression of said binding member, and recovering the binding member.

14. A specific binding member according to any one of claims 1 to 11 for use in a method of treatment or diagnosis of the human or animal body.

15. A method of preparing a specific binding member capable of binding an intracellular antigen, which method comprises:

a) providing a starting repertoire of nucleic acids encoding a VH domain which lack a CDR3 encoding

region;

b) combining said repertoire with a donor nucleic acid encoding an amino acid sequence substantially as set out as residues 99 to 106 of SEQ ID NO:1 such that said donor nucleic acid is inserted into the missing CDR3 region, so as to provide a product repertoire of nucleic acids encoding a VH domain;

c) expressing the nucleic acids of said product repertoire; and

d) selecting a specific binding member which has a maximum tumour:blood localisation ratio in a test animal of > 3:1 and optionally at said ratio an organ to blood ratio of < 1:1; and

e) recovering said binding member or the nucleic acid encoding it.

16. A method of treatment of a tumour in a human patient which comprises administering to said patient an effective amount of a specific binding member as defined in any one of claims 1 to 11.

Fig.1.

NHS76VH* Translated Sequence

Sequence Range: 1 to 351

```
         10         20         30         40         50         60
CAGGTGCAGCTGCAGGAGTCCGGCCCAGGACTGGTGAAGCCTTCGGAGACCCTGTCCCTC
GTCCACGTCGACGTCCTCAGGCCGGGTCCTGACCACTTCGGAAGCCTCTGGGACAGGGAG
 Q  V  Q  L  Q  E  S  G  P  G  L  V  K  P  S  E  T  L  S  L>          20
_____________TRANSLATION OF NHS76VH* [A]_____________>

         70         80         90        100        110        120
ACCTGCGCTGTCTCTGGTTACTCCATCAGCAGTGGTTACTACTGGGGCTGGATTCGGCAG
TGGACGCGACAGAGACCAATGAGGTAGTCGTCACCAATGATGACCCCGACCTAAGCCGTC
 T  C  A  V  S  G  Y  S  I  S [S  G  Y  Y  W  G] W  I  R  Q>  CDR1  40
_____________TRANSLATION OF NHS76VH* [A]_____________>

        130        140        150        160        170        180
CCCCCAGGGAAGGGGCTGGAGTGGATTGGGAGTATCTATCATAGTGGGAGCACCTACTAC
GGGGGTCCCTTCCCCGACCTCACCTAACCCTCATAGATAGTATCACCCTCGTGGATGATG
 P  P  G  K  G  L  E  W  I  G [S  I  Y  H  S  G  S  T  Y  Y>  CDR2  60
_____________TRANSLATION OF NHS76VH* [A]_____________>

        190        200        210        220        230        240
AACCCGTCCCTCAAGAGTCGAGTCACCATATCAGTAGACACGTCCAAGAACCAGTTCTCC
TTGGGCAGGGAGTTCTCAGCTCAGTGGTATAGTCATCTGTGCAGGTTCTTGGTCAAGAGG
[N  P  S  L  K  S] R  V  T  I  S  V  D  T  S  K  N  Q  F  S>          80
_____________TRANSLATION OF NHS76VH* [A]_____________>

        250        260        270        280        290        300
CTGAAGCTGAGCTCTGTGACCGCCGCAGACACGGCCGTGTATTACTGTGCAAGAGGGAAG
GACTTCGACTCGAGACACTGGCGGCGTCTGTGCCGGCACATAATGACACGTTCTCCCTTC
 L  K  L  S  S  V  T  A  A  D  T  A  V  Y  Y  C  A  R [G  K>  CDR3  100
_____________TRANSLATION OF NHS76VH* [A]_____________>

        310        320        330        340        350
TGGTCGAAGTTTGACTATTGGGGCCAAGGCACCCTGGTCACCGTCTCTTCA
ACCAGCTTCAAACTGATAACCCCGGTTCCGTGGGACCAGTGGCAGAGAAGT
[W  S  K  F  D  Y] W  G  Q  G  T  L  V  T  V  S  S>
_____________TRANSLATION OF NHS76VH* [A]_____________>
```

Fig.2.

NHS76VL* Translated Sequence

Sequence Range: 1 to 324

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        10         20         30         40         50         60
TCCTCTGAGCTGACTCAGGACCCTGCTGTGTCTGTGGCCTTGGGACAGACAGTCAGGATC
AGGAGACTCGACTGAGTCCTGGGACGACACAGACACCGGAACCCTGTCTGTCAGTCCTAG
 S  S  E  L  T  Q  D  P  A  V  S  V  A  L  G  Q  T  V  R  I>          20
_________________TRANSLATION OF NHS76VL* [A]_____________________>

        70         80         90        100        110        120
ACATGCCAAGGAGACAGCCTCAGAAGCTATTATGCAAGCTGGTACCAGCAGAAGCCAGGA
TGTACGGTTCCTCTGTCGGAGTCTTCGATAATACGTTCGACCATGGTCGTCTTCGGTCCT
 T  C [Q  G  D  S  L  R  S  Y  Y  A  S] W  Y  Q  Q  K  P  G>  CDR1  40
_________________TRANSLATION OF NHS76VL* [A]_____________________>

       130        140        150        160        170        180
CAGGCCCCTGTACTTGTCATCTATGGTAAAAACAACCGGCCCTCAGGGATTCCAGACCGA
GTCCGGGGACATGAACAGTAGATACCATTTTTGTTGGCCGGGAGTCCCTAAGGTCTGGCT
 Q  A  P  V  L  V  I  Y [G  K  N  N  R  P  S] G  I  P  D  R>  CDR2  60
_________________TRANSLATION OF NHS76VL* [A]_____________________>

       190        200        210        220        230        240
TTCTCTGGCTCCAGCTCAGGAAACACAGCTTCCTTGACCATCACTGGGGCTCAGGCGGAA
AAGAGACCGAGGTCGAGTCCTTTGTGTCGAAGGAACTGGTAGTGACCCCGAGTCCGCCTT
 F  S  G  S  S  S  G  N  T  A  S  L  T  I  T  G  A  Q  A  E>          80
_________________TRANSLATION OF NHS76VL* [A]_____________________>

       250        260        270        280        290        300
GATGAGGCTGACTATTACTGTAACTCCCGGGACAGCAGTGGTAACCATGTGGTATTCGGC
CTACTCCGACTGATAATGACATTGAGGGCCCTGTCGTCACCATTGGTACACCATAAGCCG
 D  E  A  D  Y  Y  C [N  S  R  D  S  S  G  N  H  V  V] F  G>  CDR3  100
_________________TRANSLATION OF NHS76VL* [A]_____________________>

       310        320
GGAGGGACCAAGCTGACCGTCCTA
CCTCCCTGGTTCGACTGGCAGGAT
 G  G  T  K  L  T  V  L>
___TRANSLATION OF NH_____>
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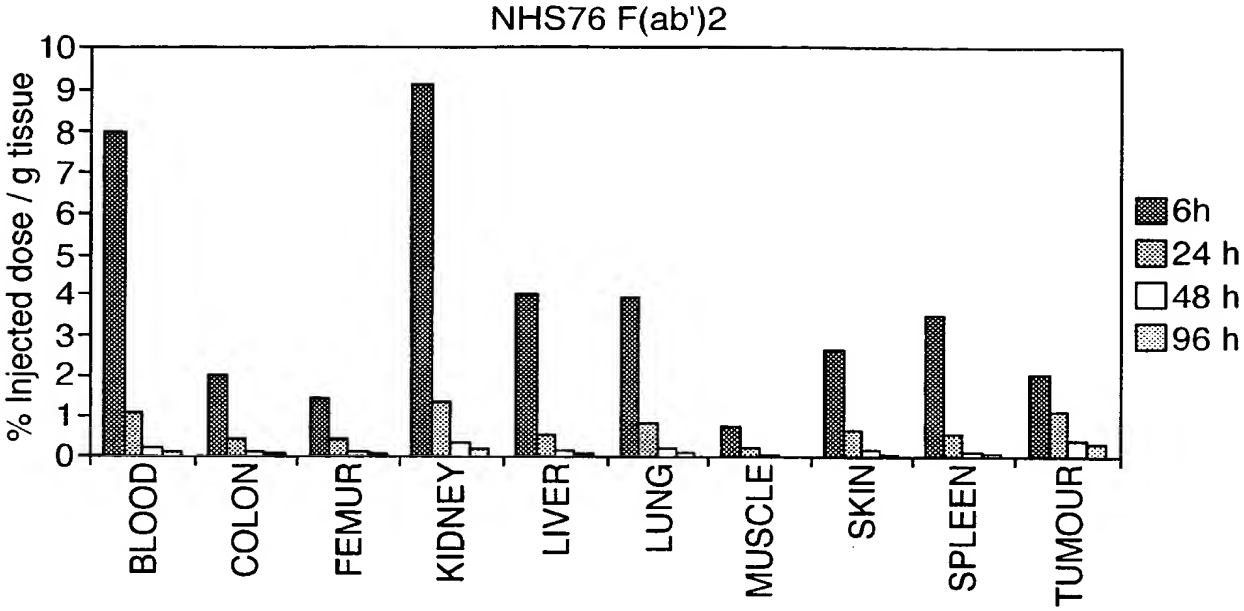
Fig.3A.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% Injected dose / g tissue
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9
8
7
6
5
4
3
2
1
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.3B.

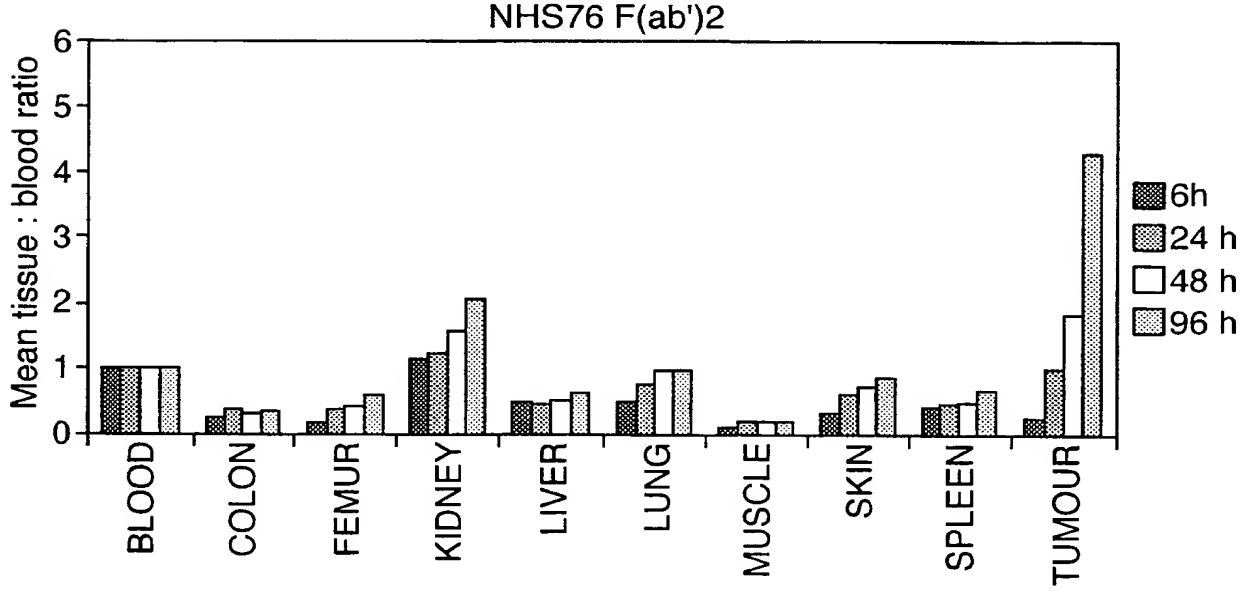
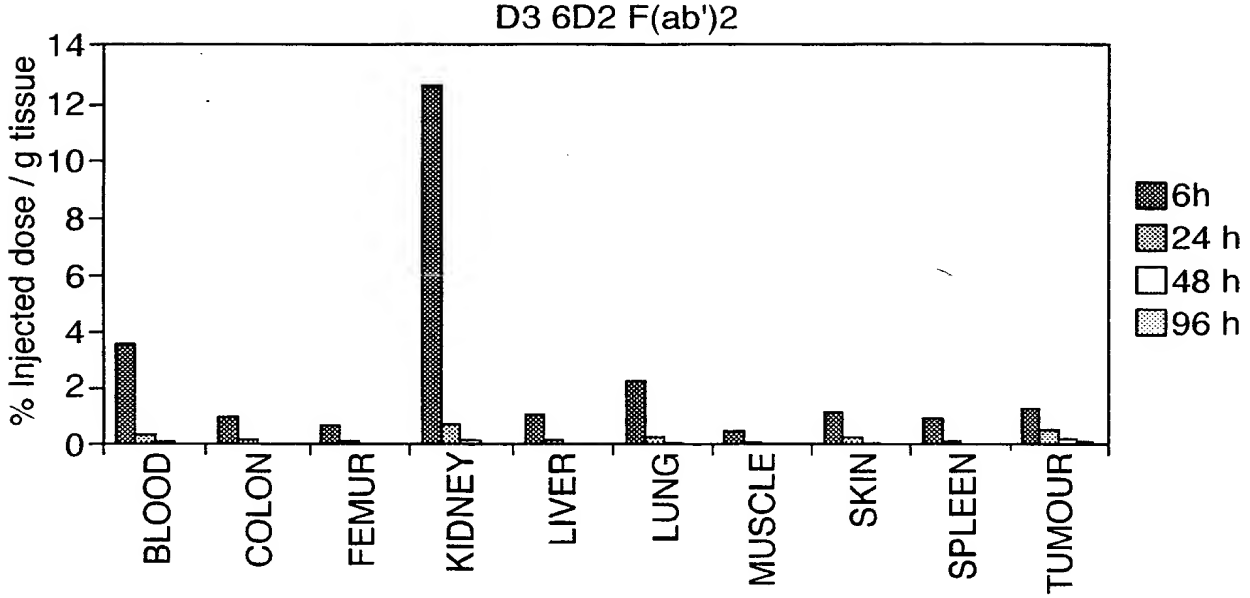

NHS76 F(ab')2
Mean tissue : blood ratio
6
5
4
3
2
1
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.4A.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% Injected dose / g tissue
14
12
10
8
6
4
2
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.4B.

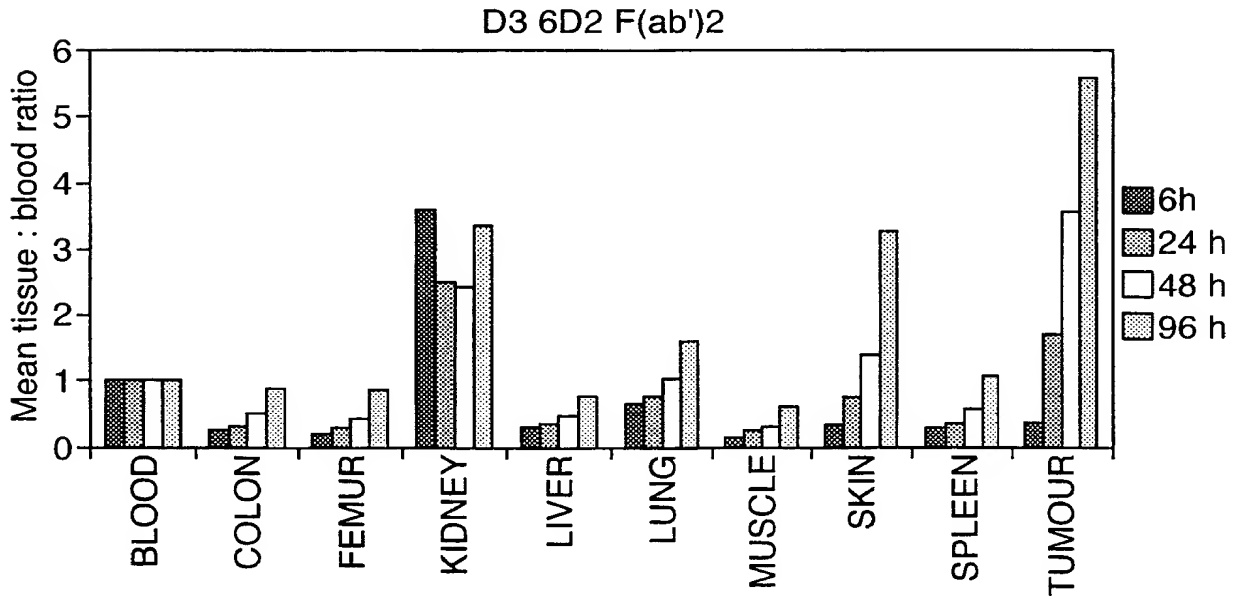
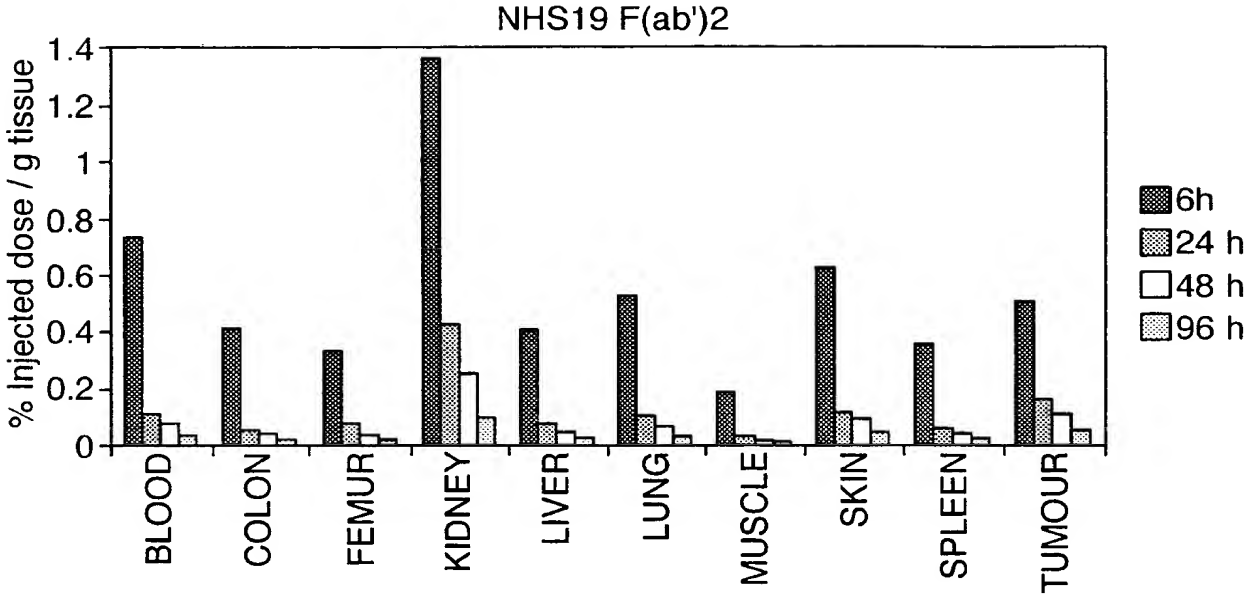

D3 6D2 F(ab')2
Mean tissue : blood ratio
6
5
4
3
2
1
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.5A.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% Injected dose / g tissue
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1.2
1
0.8
0.6
0.4
0.2
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.5B.

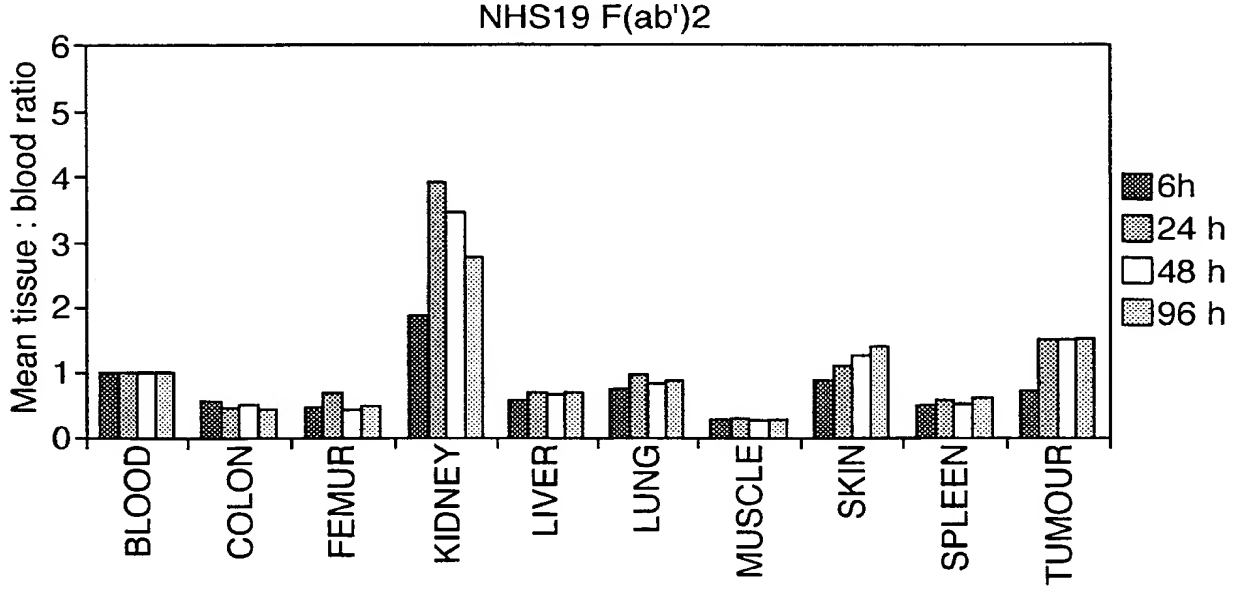

NHS19 F(ab')2
Mean tissue : blood ratio
6
5
4
3
2
1
0
BLOOD
COLON
FEMUR
KIDNEY
LIVER
LUNG
MUSCLE
SKIN
SPLEEN
TUMOUR
6h
24 h
48 h
96 h

Fig.6.

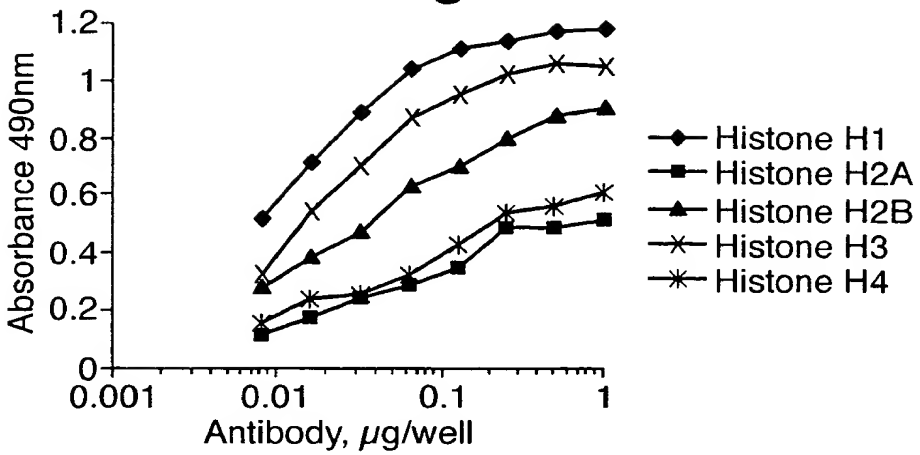
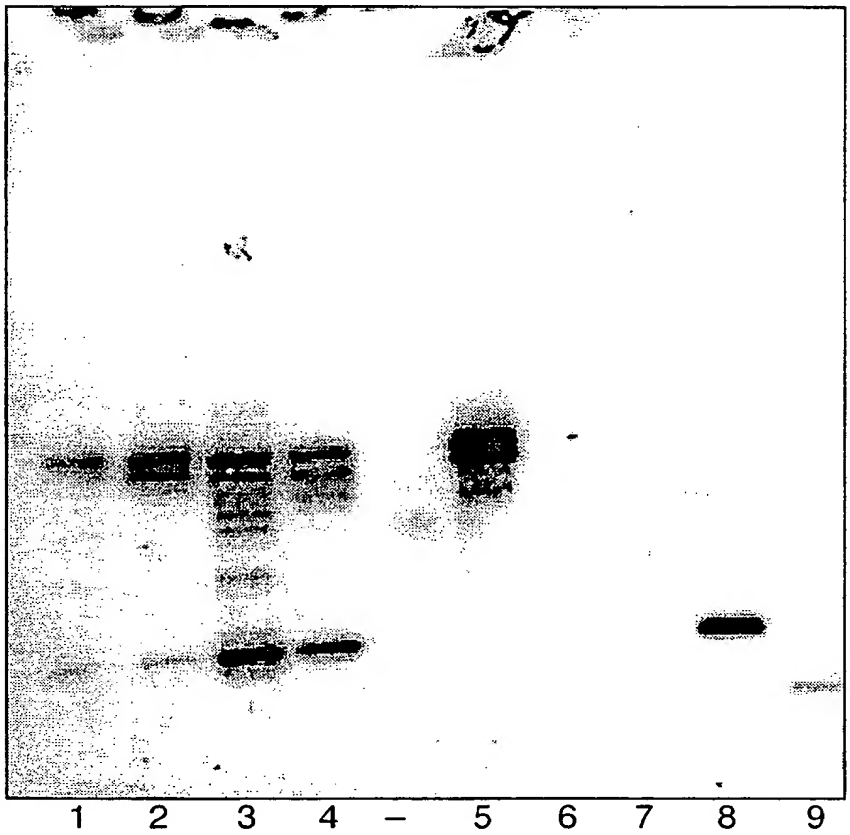


Fig.7A.



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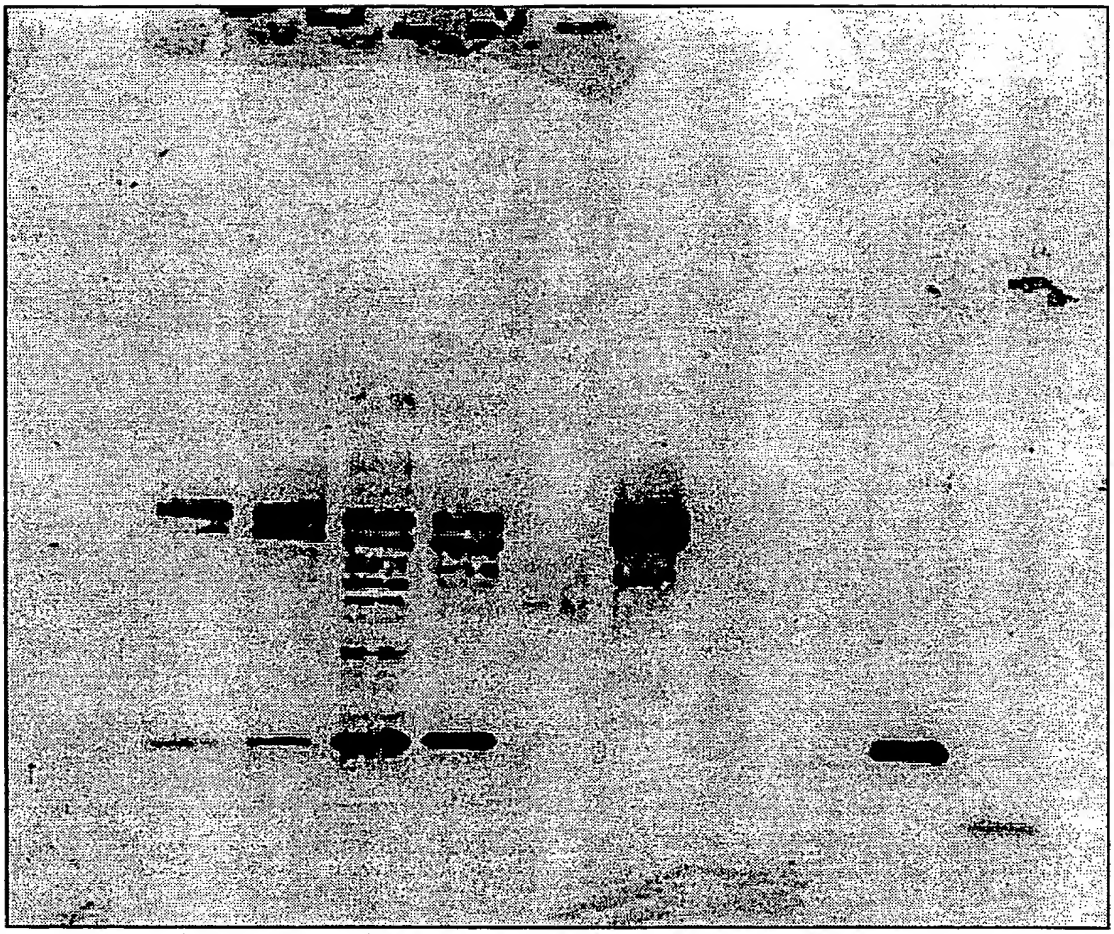
2 NSO (mouse)
3 JTC-19 (rat)
4 JH4 (guinea pig)
5 Histone H1 (bovine)
6 Histone H2A (bovine)
7 Histone H2B (bovine)
8 Histone H3 (bovine)
9 Histone H4 (bovine)

Fig.7B.



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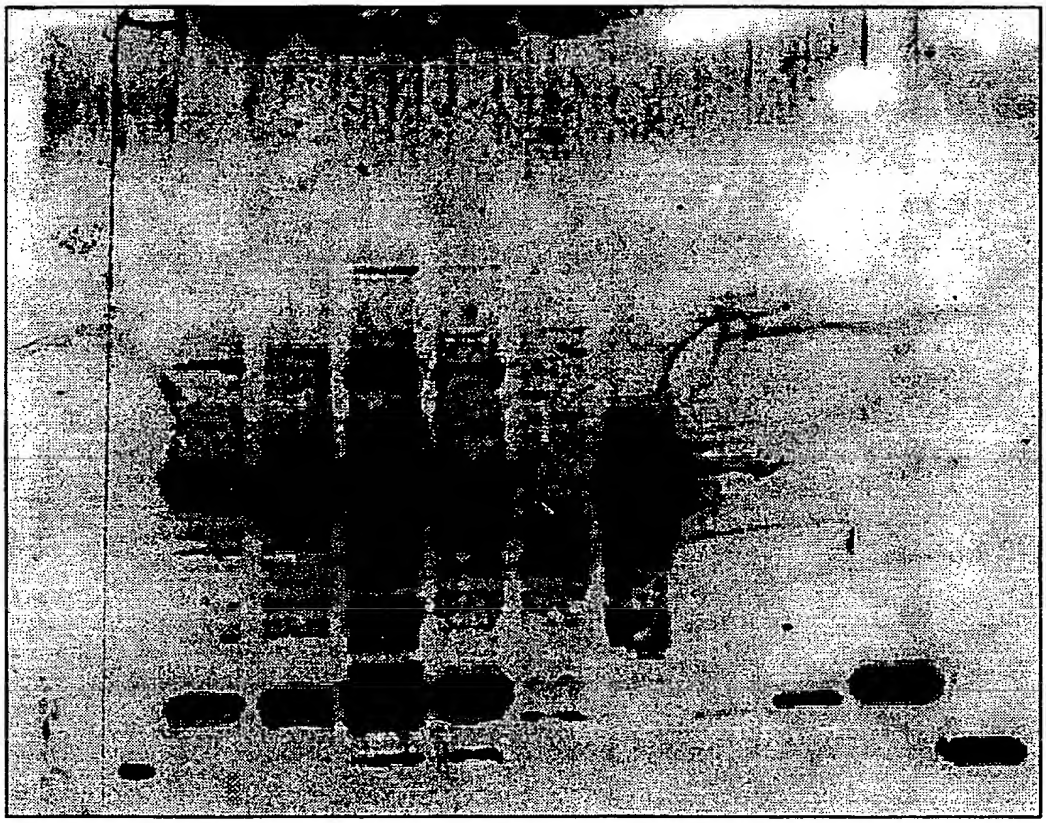
3 JTC-19 (rat)
4 JH4 (guinea pig)
5 Histone H1 (bovine)
6 Histone H2A (bovine)
7 Histone H2B (bovine)
8 Histone H3 (bovine)
9 Histone H4 (bovine)

Fig.7C.



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2 NSO (mouse)
3 JTC-19 (rat)
4 JH4 (guinea pig)
5 Histone H1 (bovine)
6 Histone H2A (bovine)
7 Histone H2B (bovine)
8 Histone H3 (bovine)
9 Histone H4 (bovine)

SEQUENCE LISTING

<110> Cambridge Antibody Technology Limited
Williams, Andrew J
Tempest, Philip R
Holtet, Thor L
Main, Sarah H
Jackson, Helen
Daramola, Olalekan

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tac tac tgg ggc tgg att cgg cag ccc cca ggg aag ggg ctg gag tgg   144
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         35                  40                  45

att ggg agt atc tat cat agt ggg agc acc tac tac aac ccg tcc ctc   192
Ile Gly Ser Ile Tyr His Ser Gly Ser Thr Tyr Tyr Asn Pro Ser Leu
     50                  55                  60

aag agt cga gtc acc ata tca gta gac acg tcc aag aac cag ttc tcc   240
Lys Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gln Phe Ser
 65                  70                  75                  80

ctg aag ctg agc tct gtg acc gcc gca gac acg gcc gtg tat tac tgt   288
Leu Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys
                 85                  90                  95

gca aga ggg aag tgg tcg aag ttt gac tat tgg ggc caa ggc acc ctg   336
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Thr Val Arg Ile Thr Cys Gln Gly Asp Ser Leu Arg Ser Tyr Tyr Ala
20 25 30

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          35                  40                  45

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 65                  70                  75                  80

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            20                  25                  30

Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
          35                  40                  45

Gly Lys Asn Asn Arg Pro Ser Gly Ile Pro Asp Arg Phe Ser Gly Ser
     50                  55                  60
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<210> 6
<211> 21
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<220>
<223> Description of Artificial Sequence: Primer

<400> 6
gtcgtctttc cagacgttag t 21

<210> 7
<211> 23
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<220>
<223> Description of Artificial Sequence: Primer

<400> 7
agcggataac aatttcacac agg 23

<210> 8
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<220>
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<400> 8
ctcttctgag atgagttttt g 21

<210> 9
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<220>
<223> Description of Artificial Sequence: Primer

<400> 9
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<210> 10
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<220>
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<400> 10
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<210> 11
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<220>
<223> Description of Artificial Sequence: Primer

<400> 11
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<210> 12
<211> 35
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<220>
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<400> 12
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<210> 13
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<400> 13
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<210> 14
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<400> 14
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<210> 15
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<220>
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<210> 16
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<212> DNA
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<220>
<223> Description of Artificial Sequence: Primer

<400> 16
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<210> 17
<211> 25
<212> DNA
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<220>
<223> Description of Artificial Sequence: Primer

<400> 17
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<210> 18
<211> 25
<212> DNA
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<220>
<223> Description of Artificial Sequence: Primer

<400> 18
gccctcaggg attccagacc gattc 25

<210> 19
<211> 25
<212> DNA
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<220>
<223> Description of Artificial Sequence: Primer

<400> 19
gaatcggtct ggaatccctg agggc 25

<210> 20
<211> 53
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: Primer

<400> 20
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<210> 21
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<212> DNA
<213> Homo sapiens

<400> 21
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gttcttggac gtgtctactg atatggtgac tcgactcttg agggacgggt tgtagtaggt 180
gctcccacta tgatagatac tcccaatcca ctccagcccc ttccctgggg gctgccgaat 240
ccagccccag tagtaaccac tgctgatgga gtaaccagag acagcgcagg tgagggacag 300
ggtctccgaa ggcttcacca gtcctgggcc ggactcctgc agctgcacct g 351

<210> 22
<211> 324
<212> DNA
<213> Homo sapiens

<400> 22
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gtttcctgag ctggagccag agaatcggtc tggaatccct gagggccggt tgtttttacc 180
atagatgaca agtacagggg cctgtcctgg cttctgctgg taccagcttg cataatagct 240
tctgaggctg tctccttggc atgtgatcct gactgtctgt cccaaggcca cagacacagc 300
agggtcctga gtcagctcag agga 324